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(71) Applicant: WILLIAM BEAUMONT HOSPITAL
[US/US]; 3601 West Thirteen Mile Road, Royal Oak, MI
48073-6769 (US).

(72) Inventors: JAFFREY, David, A.; 2476 Lincoln Road,
Windsor, Ontario N8W 2R7 (CA). WONG, John, W.; 726
Tennyson Downs Court, Bloomfield Hills, MI 48304 (US).
SIEWERDSEN, Jeffrey, H.; 2383 Timbercrest Court,
Ann Arbor, MI 48105 (US).

(74) Agent: FREEMAN, John, C.; Brinks Hofer Gilson &
Lione, P.O. Box 10087, Chicago, IL 60610 (US).

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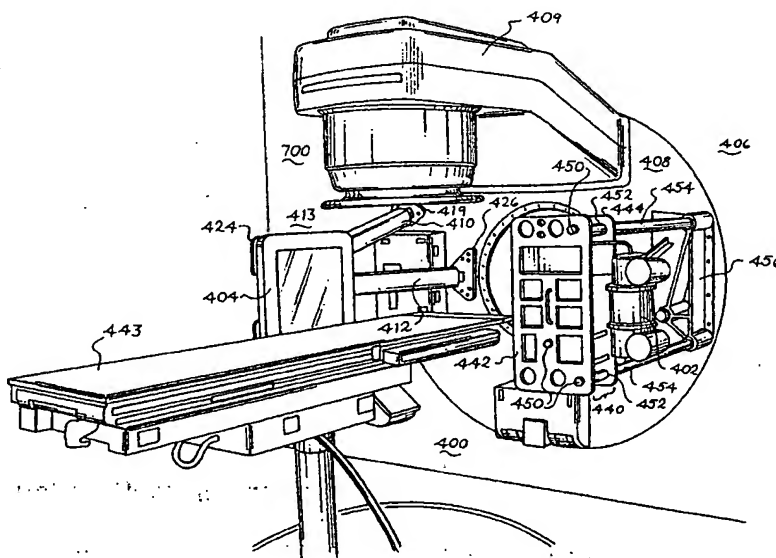
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(54) Title: CONE-BEAM COMPUTERIZED TOMOGRAPHY WITH A FLAT-PANEL IMAGER



(57) Abstract: A radiation therapy system that includes a radiation source that moves about a path and directs a beam of radiation towards an object and a cone-beam computer tomography system. The cone-beam computer tomography system includes an x-ray source that emits an x-ray beam in a cone-beam form towards an object to be imaged and an amorphous silicon flat-panel imager receiving x-rays after they pass through the object, the imager providing an image of the object. A computer is connected to the radiation source and the cone beam computerized tomography system, wherein the computer receives the image of the object and based on the image sends a signal to the radiation source that controls the path of the radiation source.

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CONE-BEAM COMPUTERIZED TOMOGRAPHY WITH A FLAT-PANEL IMAGER

Applicants claim, under 35 U.S.C. § 119(e), the benefit of priority of the filing date of February 18, 2000, of U.S. Provisional Patent Application Serial Number 60/183,590, filed on the aforementioned date, the entire contents of which are incorporated herein by reference.

BACKGROUND OF THE INVENTION

Field of the Invention

The present invention relates generally to a cone-beam computed tomography system and, more particularly, to a cone-beam computed tomography system that employs an amorphous silicon flat-panel imager for use in radiotherapy applications where the images of the patient are acquired with the patient in the treatment position on the treatment table.

Discussion of the Related Art

Radiotherapy involves delivering a prescribed tumorcidal radiation dose to a specific geometrically defined target or target volume. Typically, this treatment is delivered to a patient in one or more therapy sessions (termed fractions). It is not uncommon for a treatment schedule to involve twenty to forty fractions, with five fractions delivered per week. While radiotherapy has proven successful in managing various types and stages of cancer, the potential exists for increased tumor control through increased dose. Unfortunately, delivery of increased dose is limited by the presence of adjacent normal structures and the precision of beam delivery. In some sites, the diseased target is directly adjacent to radiosensitive normal structures. For example, in the treatment of prostate cancer, the prostate and rectum are directly adjacent. In this situation, the prostate is the targeted volume and the maximum deliverable dose is limited by the wall of the rectum.

In order to reduce the dosage encountered by radiosensitive normal structures, the location of the target volume relative to the radiation therapy source must be known precisely in each treatment session in order to accurately deliver a tumorcidal dose while minimizing complications in normal tissues. Traditionally, a radiation therapy treatment plan is formed based on the location and orientation of the lesion and surrounding structures in an initial computerized tomography or magnetic resonance image. However, the location and orientation of the lesion may vary during the course of treatment from that used to form the radiation therapy treatment plan. For example, in each treatment session, systematic and/or random variations in patient setup (termed interfraction setup errors) and in the location of the lesion relative to surrounding anatomy (termed interfraction organ motion errors) can each change the location and orientation of the lesion at the time of treatment compared to that assumed in the radiation therapy treatment plan. Furthermore, the location and orientation of the lesion can vary during a single treatment session (resulting in intrafraction errors) due to normal biological processes, such as breathing, peristalsis, etc. In the case of radiation treatment of a patient's prostate, it is necessary to irradiate a volume that is enlarged by a margin to guarantee that the prostate always receives a prescribed dose due to uncertainties in patient positioning and daily movement of the prostate within the patient. Significant dose escalation may be possible if these uncertainties could be reduced from current levels (~10mm) to 2-3mm.

Applying large margins necessarily increases the volume of normal tissue that is irradiated, thereby limiting the maximum dose that can be delivered to the lesion without resulting in complication in normal structures. There is strong reason to believe that increasing the dose delivered to the lesion can result in more efficacious treatment. However, it is often the case that the maximum dose that can be safely delivered to the target volume is limited by the associated dose to surrounding normal structures incurred through the use of margins. Therefore, if one's knowledge of the location and orientation of the lesion at the time of treatment can be increased, then

margins can be reduced, and the dose to the target volume can be increased without increasing the risk of complication in normal tissues.

A number of techniques have been developed to reduce uncertainty associated with systematic and/or random variations in lesion location resulting from interfraction and intrafraction errors. These include patient immobilization techniques (e.g., masks, body casts, bite blocks, etc.), off-line review processes (e.g., weekly port films, population-based or individual-based statistical approaches, repeat computerized tomography scans, etc.), and on-line correction strategies (e.g., pre-ports, MV or kV radiographic or fluoroscopic monitoring, video monitoring, etc.).

It is believed that the optimum methodology for reducing uncertainties associated with systematic and/or random variations in lesion location can only be achieved through using an on-line correction strategy that involves employing both on-line imaging and guidance system capable of detecting the target volume, such as the prostate, and surrounding structures with high spatial accuracy.

An on-line imaging system providing suitable guidance has several requirements if it is to be applied in radiotherapy of this type. These requirements include contrast sensitivity sufficient to discern soft-tissue; high spatial resolution and low geometric distortion for precise localization of soft-tissue boundaries; operation within the environment of a radiation treatment machine; large field-of-view (FOV) capable of imaging patients up to 40cm in diameter; rapid image acquisition (within a few minutes); negligible harm to the patient from the imaging procedure (e.g., dose much less than the treatment dose); and compatibility with integration into an external beam radiotherapy treatment machine.

Several examples of known on-line imaging systems are described below. For example, strategies employing x-ray projections of the patient (e.g., film, electronic portal imaging devices, kV radiography/fluoroscopy, etc.) typically show only the location of bony anatomy and not soft-tissue structures. Hence, the location of a soft-tissue target volume must be inferred from the location of bony landmarks. This obvious shortcoming can be

alleviated by implanting radio-opaque markers on the lesion; however, this technique is invasive and is not applicable to all treatment sites. Tomographic imaging modalities (e.g., computerized tomography, magnetic resonance, and ultrasound), on the other hand, can provide information regarding the location of soft-tissue target volumes. Acquiring computerized tomography images at the time of treatment is possible, for example, by incorporating a computerized tomography scanner into the radiation therapy environment (e.g., with the treatment table translated between the computerized tomography scanner gantry and the radiation therapy gantry along rails) or by modifying the treatment machine to allow computerized tomography scanning. The former approach is a fairly expensive solution, requiring the installation of a dedicated computerized tomography scanner in the treatment room. The latter approach is possible, for example, by modifying a computer tomography scanner gantry to include mechanisms for radiation treatment delivery, as in systems for tomotherapy. Finally, soft-tissue visualization of the target volume can in some instances be accomplished by means of an ultrasound imaging system attached in a well-defined geometry to the radiation therapy machine. Although this approach is not applicable to all treatment sites, it is fairly cost-effective and has been used to illustrate the benefit of on-line therapy guidance.

As illustrated in FIGS. 1(a)-(c), a typical radiation therapy system 100 incorporates a 4-25 MV medical linear accelerator 102, a collimator 104 for collimating and shaping the radiation field 106 that is directed onto a patient 108 who is supported on a treatment table 110 in a given treatment position. Treatment involves irradiation of a lesion 112 located within a target volume with a radiation beam 114 directed at the lesion from one or more angles about the patient 108. An imaging device 116 may be employed to image the radiation field 118 transmitted through the patient 108 during treatment. The imaging device 116 for imaging the radiation field 118 can be used to verify patient setup prior to treatment and/or to record images of the actual radiation fields delivered during treatment. Typically, such images suffer from poor

contrast resolution and provide, at most, visualization of bony landmarks relative to the field edges.

Another example of a known on-line imaging system used for reducing uncertainties associated with systematic and/or random variations in lesion location is an X-ray cone-beam computerized tomography system. Mechanical operation of a cone beam computerized tomography system is similar to that of a conventional computerized tomography system, with the exception that an entire volumetric image is acquired through a single rotation of the source and detector. This is made possible by the use of a two-dimensional (2-D) detector, as opposed to the 1-D detectors used in conventional computerized tomography. There are constraints associated with image reconstruction under a cone-beam geometry. However, these constraints can typically be addressed through innovative source and detector trajectories that are well known to one of ordinary skill in the art.

As mentioned above, a cone beam computerized tomography system reconstructs three-dimensional (3-D) images from a plurality of two-dimensional (2-D) projection images acquired at various angles about the subject. The method by which the 3-D image is reconstructed from the 2-D projections is distinct from the method employed in conventional computerized tomography systems. In conventional computerized tomography systems, one or more 2-D slices are reconstructed from one-dimensional (1-D) projections of the patient, and these slices may be "stacked" to form a 3-D image of the patient. In cone beam computerized tomography, a fully 3-D image is reconstructed from a plurality of 2-D projections. Cone beam computerized tomography offers a number of advantageous characteristics, including: formation of a 3-D image of the patient from a single rotation about the patient (whereas conventional computerized tomography typically requires a rotation for each slice); spatial resolution that is largely isotropic (whereas in conventional computerized tomography the spatial resolution in the longitudinal direction is typically limited by slice thickness); and considerable flexibility in the imaging geometry. Such technology has been employed in applications such as

micro-computerized tomography, for example, using a kV x-ray tube and an x-ray image intensifier tube to acquire 2-D projections as the object to be imaged is rotated, e.g., through 180° or 360°. Furthermore, cone beam computerized tomography has been used successfully in medical applications such as computerized tomography angiography, using a kV x-ray tube and an x-ray image intensifier tube mounted on a rotating C-arm.

The development of a kV cone-beam computerized tomography imaging system for on-line tomographic guidance has been reported. The system consists of a kV x-ray tube and a radiographic detector mounted on the gantry of a medical linear accelerator. The imaging detector is based on a low-noise charge-coupled device (CCD) optically coupled to a phosphor screen. The poor optical coupling efficiency ($\sim 10^{-4}$) between the phosphor and the CCD significantly reduces the detective quantum efficiency (DQE) of the system. While this system is capable of producing cone beam computerized tomography images of sufficient quality to visualize soft tissues relevant to radiotherapy of the prostate, the low DQE requires imaging doses that are a factor of 3-4 times larger than would be required for a system with an efficient coupling (e.g. $\sim 50\%$ or better) between the screen and detector.

Another example of a known auxiliary cone beam computerized tomography imaging system is shown in FIG. 2. The auxiliary cone beam computerized tomography imaging system 200 replaces the CCD-based imager of FIGS. 1(a)-(c) with a flat-panel imager. In particular, the imaging system 200 consists of a kilovoltage x-ray tube 202 and a flat panel imager 204 having an array of amorphous silicon detectors that are incorporated into the geometry of a radiation therapy delivery system 206 that includes an MV x-ray source 208. A second flat panel imager 210 may optionally be used in the radiation therapy delivery system 206. Such an imaging system 200 could provide projection radiographs and/or continuous fluoroscopy of the lesion 212 within the target volume as the patient 214 lies on the treatment table 216 in the treatment position. If the geometry of the imaging system 200 relative to the system 206 is known, then the resulting kV projection images could be used to modify patient setup and improve somewhat the precision of radiation

treatment. However, such a system 200 still would not likely provide adequate visualization of soft-tissue structures and hence be limited in the degree to which it could reduce errors resulting from organ motion.

Accordingly, it is an object of the present invention to generate KV projection images in a cone beam computerized tomography system that provide adequate visualization of soft-tissue structures so as to reduce errors in radiation treatment resulting from organ motion.

BRIEF SUMMARY OF THE INVENTION

One aspect of the present invention regards a radiation therapy system that includes a radiation source that moves about a path and directs a beam of radiation towards an object and a cone-beam computer tomography system. The cone-beam computer tomography system includes an x-ray source that emits an x-ray beam in a cone-beam form towards an object to be imaged and an amorphous silicon flat-panel imager receiving x-rays after they pass through the object, the imager providing an image of the object. A computer is connected to the radiation source and the cone beam computerized tomography system, wherein the computer receives the image of the object and based on the image sends a signal to the radiation source that controls the path of the radiation source.

A second aspect of the present invention regards a method of treating an object with radiation that includes moving a radiation source about a path, directing a beam of radiation from the radiation source towards an object and emitting an x-ray beam in a cone beam form towards the object. The method further includes detecting x-rays that pass through the object due to the emitting an x-ray beam with an amorphous silicon flat-panel imager, generating an image of the object from the detected x-rays and controlling the path of the radiation source based on the image.

Each aspect of the present invention provides the advantage of generating KV projection images in a cone beam computerized tomography system that provide adequate visualization of soft-tissue structures so as to reduce errors in radiation treatment resulting from organ motion.

Each aspect of the present invention provides an apparatus and method for improving the precision of radiation therapy by incorporating a cone beam computerized tomography imaging system in the treatment room, the 3-D images from which are used to modify current and subsequent treatment plans.

Each aspect of the present invention represents a significant shift in the practice of radiation therapy. Not only does the high-precision, image-guided system for radiation therapy address the immediate need to improve the probability of cure through dose escalation, but it also provides opportunity for broad innovation in clinical practice.

Each aspect of the present invention may permit alternative fractionation schemes, permitting shorter courses of therapy and allowing improved integration in adjuvant therapy models.

Each aspect of the present invention provides valuable imaging information for directing radiation therapy also provides an explicit 3-D record of intervention against which the success or failure of treatment can be evaluated, offering new insight into the means by which disease is managed.

Additional objects, advantages and features of the present invention will become apparent from the following description and the appended claims when taken in conjunction with the accompanying drawings.

BRIEF DESCRIPTION OF THE DRAWINGS

FIGS. 1(a)-(c) schematically show the geometry and operation of a conventional radiation therapy apparatus;

FIG. 2 schematically shows a perspective view of a known radiation therapy apparatus including an auxiliary apparatus for cone beam computerized tomography imaging;

FIG. 3 is a diagrammatic view of a bench-top cone beam computerized tomography system employing a flat-panel imager, according to a first embodiment of the present invention;

FIG. 4 is a schematic illustration of the geometry and procedures of the cone beam computerized tomography system shown in FIG. 3;

FIGS. 5(a) - 5(d) are graphs depicting the fundamental performance characteristics of the flat-panel imager used in the cone beam computerized tomography system of FIG. 3;

5 FIGS. 6(a) - 6(d) show various objects used in tests to investigate the performance of the cone beam computerized tomography system of the present invention, including a uniform water cylinder, six low-contrast inserts in a water bath, a steel wire under tension with a water bath, and an euthanized rat, respectively;

10 FIGS. 7(a) - 7(d) depict uniformity of response of the cone beam computerized tomography system of the present invention, including axial and sagittal slices through volume images of a uniform water bath, radial profiles, and a vertical signal profile, respectively;

15 FIGS. 8(a) - 8(d) illustrate the noise characteristics of the cone beam computerized tomography system of the present invention, including axial and sagittal noise images from volume reconstructions of a uniform water bath, radial noise profiles, and vertical noise profiles, respectively;

FIGS. 9(a) - 9(b) depict response linearity and voxel noise, respectively, for the cone beam computerized tomography system of the present invention and a conventional computerized tomography scanner;

20 FIGS. 10(a) - 10(c) depict the noise-power spectrum from the cone beam computerized tomography system of the present invention, including a gray scale plot of the axial noise-power spectrum, the noise-power spectrum measured at various exposures, and the noise-power spectrum for the cone beam computerized tomography system compared to a conventional
25 computerized tomography scanner, respectively;

FIGS. 11 (a) - 11 (b) depict the spatial resolution of the cone beam computerized tomography system of the present invention, including the surface plot of an axial slice image of the thin steel wire shown in FIG. 6(c) and the modulation transfer function measured for the cone beam
30 computerized tomography system and for a conventional computerized tomography scanner, respectively;

FIGS. 12(a) - 12(b) show images of a low-contrast phantom obtained from the cone beam computerized tomography system of the present invention and a conventional computerized tomography scanner, respectively;

5 FIGS. 13(a) - 13(i) show cone beam computerized tomography images of the euthanized rat shown in FIG. 6(d), including regions of the lungs (FIGS. 13(a) - 13(c)), the kidneys (FIGS. 13(d)-13(f)), and the lower spine (FIGS. 13(g) - 13(i));

10 FIGS. 14(a) - 14(d) show volume renderings of cone beam computerized tomography images of the euthanized rat shown in FIG. 6(d) illustrating the degree of spatial resolution achieved in delineating structures of the vertebra, including volume renderings with axial and sagittal cut planes showing the skeletal anatomy along with soft-tissue structures of the abdomen, volume renderings with axial and sagittal cut planes, window to show skeletal features only, a magnified view of a region of the spine and ribs
15 of the rat, and a magnified view of a part of two vertebra, respectively;

FIGS. 15(a) - 15(b) depict the axial images of euthanized rat shown in FIG. 6(d) obtained from the cone beam computerized tomography system of the present invention and a conventional computerized tomography scanner, respectively;

20 FIG. 16 is a graph showing detected quantum efficiency calculated as a function of exposure for an existing and hypothetical flat-panel imager configuration;

FIGS. 17(a)-(e) are diagrammatic views of several angular orientations of a wall-mounted cone beam computerized tomography system employing a flat-panel imager, according to a second embodiment of the present invention;
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FIG. 18 shows a side view of the cone beam computerized tomography system of FIG. 17 when employing a first embodiment of a support for a flat-panel imager according to the present invention;

30 FIG. 19(a) shows a perspective exploded view of a mounting to be used with the support for a flat-panel imager of FIG. 18;

FIG. 19(b) shows a perspective exploded view of a rotational coupling to be used with the mounting of FIG. 19(a);

FIGS. 20(a)-(b) schematically shows a front view of the wall-mounted cone beam computerized tomography system of FIG. 17 when employing a second embodiment of a support for a flat-panel imager according to the present invention;

5 FIGS. 21(a)-(b) schematically shows a front view of the wall-mounted cone beam computerized tomography system of FIG. 17 when employing a third embodiment of a support for a flat-panel imager according to the present invention;

10 FIG. 22 is a diagrammatic view of a portable cone beam computerized tomography system employing a flat-panel imager according to fifth embodiment of the present invention;

FIGS. 23(a)-(d) are diagrammatic sketches illustrating the geometry and operation of the cone beam computerized tomography imaging systems of FIGS. 17-22;

15 FIG. 24 is a flow-chart showing an embodiment of the processes involved in acquiring a cone beam computerized image for the cone beam computerized tomography imaging systems of FIGS. 17-22;

20 FIG. 25 is a perspective drawing illustrating an embodiment of a method for geometric calibration of the imaging and treatment delivery systems of FIGS. 17-22; and

25 FIG. 26 is a flow-chart showing an embodiment of the processes involved in the image-guided radiation therapy systems of FIGS. 17-22, based on cone beam computerized tomography imaging of a patient, on-line correction of setup errors and organ motion, and off-line modification of subsequent treatment plans.

PREFERRED EMBODIMENTS OF THE INVENTION

30 A bench-top cone beam computerized tomography (CBCT) system 300 is shown in FIG. 3, according to an embodiment of the present invention. The CBCT system 300 was constructed to mimic the geometry of the CBCT scanner currently installed on a linear accelerator, with a source-to-axis distance of 1000 mm and a source-detector distance of 1600 mm. The

primary components of the system 300 include an x-ray tube 302, a rotation stage 304 and flat-panel imager (FPI) 306. These components are rigidly mounted to an optical bench 308. The relative position of these components is controlled by three translation stages, including an xobject stage 310, a yobject stage 312 and a yimage stage 314, which are used during initial setup to accurately determine and control the imaging geometry. The cone beam computerized tomography system 300 generates images of an object 316, identified throughout as a phantom, mounted on the rotation stage 304. Each stage 310, 312 and 314 contains a home or limit switch, and the imaging geometry is referenced to the location of these switches with a reproducibility of ± 0.01 mm. The specific geometries used in the discussion herein are shown in FIG. 4, and are set to simulate the imaging geometry that would be implemented for a cone beam computerized tomography system incorporated on a radiotherapy treatment machine. Table 1 below shows the parameters of the system 300.

A set of alignment lasers 318 allow visualization of the axis of rotation 320 and the source plane perpendicular to the axis of rotation 320 and intersects focal spot 322 of the x-ray source or tube 302. The axis of rotation 320 is positioned such that it intersects the central ray 324 between the focal spot 322 and the detector plane 326 ($+0.01$ mm). The flat plane imager 326 is positioned such that the piercing point (i.e., the intersection of the central ray and the image plane) is centered on the imaging array (i.e., between columns #256 and #257, ± 0.01 mm), with a quarter-pixel offset applied to give improved view sampling for cone beam computerized tomography acquisitions in which the object 316 is rotated through 360° . The stage 310 is controlled manually by means of a positioning micrometer. The source-to-object (SOD) and source-to-image (SID) distances were measured to within ± 0.5 mm and give an objection magnification of 1.60, equal to that of the imaging system on the linear accelerator. The cone angle for this geometry is -7.1° .

Radiographic exposures used in the acquisition procedure are produced under computer control with a 300 kHU x-ray tube 302, such as

General Electric Maxi-ray 75 and a 100 kW generator, such as the General Electric MSI-800. The tube 302 has a total minimum filtration of 2.5 mm A1, with an additional filtration of 0.127 mm Cu to further harden the beam, and a nominal focal spot size of 0.6 mm. The 100 kV beam is characterized by first and second HVLs of 5.9 and 13.4 mm A1, respectively. The accelerating potential of the generator was monitored over a one-week period and was found to be stable to within $\pm 1\%$. All exposures were measured using an x-ray multimeter, such as the RTI Electronics, Model PMX-III with silicon diode detector.

The exposures for the cone beam computerized tomography acquisitions are reported in terms of exposure to air at the axis of rotation in the absence of the object 316. The same method of reporting exposure can be used for the images acquired on the conventional scanner. For the conventional scanner, the exposure per unit charge is measured with the gantry rotation disabled and the collimators set for a 10 mm slice thickness, thereby guaranteeing complete coverage of the silicon diode. The exposure per unit charge at 100 kVp was 9.9 mR/mAs and 14.9 mR/mAs for the bench-top and conventional scanners, respectively.

The flat panel imager 306 can be the EG&G Heimann Optoelectronics (RID 512-400 AO) that incorporates a 512 x 512 array of a-Si:H photodiodes and thin-film transistors. The electro-mechanical characteristics of the imager are shown in Table 1. The flat plane imager 306 is read-out at one of eight present frame rates (up to 5 frames per second) and operates asynchronously of the host computer 328 schematically shown in FIG. 4. The analog signal from each pixel is integrated by ASIC amplifiers featuring correlated double-sampling noise reduction circuitry. Digitization is performed at 16 bit resolution. The values are transferred via an RS-422 bus to a hardware buffer in the host computer 328. The processor in the host computer 328 is interrupted when a complete frame is ready for transfer to host memory.

Table 1

CBCT Characteristic	Value
Acquisition Geometry	
Source-axis-distance (S_{AD})	103.3 cm
Source-imager-distance (S_{ID})	165.0 cm
Cone angle	7.1°
Maximum angular rotation rate	0.5°/sec
Field of view (FOV)	12.8 cm
X-ray Beam/Exposure Characteristics	
Beam energy	100 kVp
Added filtration	1.5mm A1 + 0.129 mm Cu
Beam quality	$HVL_1 = 5.9 \text{ MM A1}$
	$HVL_2 = 13.4 \text{ MM A1}$
Scatter-to-primary ratio	0.18, 1:5 (11 cm object)
Frame time	6.4 sec
Tube output at (SAD)	9.34 mR/mAs
Exposure rate (at SID)	3.65 mR/mAs
Flat-Panel Imager	
Designation	RID 512-400 AO
Array format	512 x 512 pixels
Pixel pitch	400 μm

CBCT Characteristic	Value
Area	~20.5 20.5 cm ²
Pixel fill factor	0.80
Photodiode charge capacity	~62 Pc
ASIC amplifier charge capacity	~23 pC
ASIC amplifier noise	~12,700 e
ADC bit-depth	16 bit
TFT thermal noise (on)	~1800 e
Photodiode Shot Noise (1 fps)	~1200 e
Digitization noise	~630 e
Nominal frame rate	0.16 fps
Maximum frame rate	5 fps
X-ray converter	133 mg/cm ² Gd ₂ O ₂ S:Tb
Acquisition Procedure	
Number of projections	300
Angular increment	1.2°
Total rotation angle	360°
Maximum angular rotation rate	05°/s
Reconstruction Parameters	
Reconstruction matrix	561 x 561 x (1-512), 281 x 281 x (1-500)
Voxel size	0.25 x 0.25 x 0.25 mm ² , 0.5 x 0.5 x 0.25

CBCT Characteristic	Value
W. parameter	1.60
γ , cutoff frequency modification	1.0
α , modified Hamming filter parameter	0.50
Range of convolution	± 25 mm

The cone-beam scanning procedure includes a repeated sequence of radiographic exposure, array readout, and object rotation. The timing of this procedure is driven by the asynchronous frame clock of the flat plane imager readout electronics. A conservative frame time of 6.4 s was used. Between the periodic frame transfers from the flat plane imager 306, the host computer advances the motorized rotation stage 304 and triggers the x-ray generator or tube 302. The rotor of the x-ray tube 302 remains spinning throughout the scanning procedure. The control software allows the operator to specify the number of frames between exposures. This was designed as a mechanism to investigate methods of reducing the amount of lag in sequential projections. The detector signal from a group of nine pixels in the bare-beam region of the flat plane imager 306 is monitored to measure and verify the stability of each radiographic exposure. Exposures outside tolerance are trapped and repeated at the same projection angle. Each projection image is written to hard disk between frame transfer and motor rotation. After the projections are acquired, a set of flood and dark field images (20 each) are collected to construct gain and offset images for flat-field processing of the projection images.

In addition to gain and offset corrections, median filtration (3x3) is performed using a pre-constructed map of unstable pixels. Finally, the signal in each projection is normalized to account for small variations in x-ray exposure, this is performed using a cluster of nine pixels in the periphery of the detector well outside the objects shadow.

A volumetric computerized tomography data set is reconstructed from the projections using a filtered back-projection technique. The filter used in the reconstruction is constructed using Webb's three-parameter formula. The parameters and their corresponding values are shown in Table 1. In the current configuration, the reconstruction field of vision is limited to a 12.4 cm diameter cylinder, approximately 12.1 cm in length; the lateral extent of objects to be reconstructed must lie well-within this cylinder. The voxel values in the resulting volumetric data sets are scaled linearly to produce a mean CT number of zero in air and 1000 in water. The time required to filter (100 element kernel) and back-project a single projection (512 x 512) on to a 281 x 281 x 500 voxel data set was 1 minute and 21 seconds.

The basic signal and noise characteristics of the flat plane imager 306 were measured. The detector gain and linearity are presented in FIG. 5(a). For an x-ray beam energy of 120 kVp, the detector gain was measured to be 18.2×10^5 e/mR/pixel (17.8×10^6 e/mR at 100 kVp). The detector exhibits excellent linearity with exposure up to 50% of its sensitive range (5 mR). The various additive electronic noise sources and their magnitudes are listed in Table 1. The total additive electronic noise is found to depend upon frame time, ranging from 13,300 e at a frame time of 200 ms to 22,500 e at a frame time of 25.6s. The amplifier noise (12,700 e) is the dominant component at high frame rates. The significance of amplifier noise on the zero-frequency detective quantum efficiency (DQE) was studied using a cascaded system model that analyzes signal and noise propagation in the FPI 306.

FIG. 5(b) shows the dependence of detective quantum efficiency on exposure for the RID 512-400AO, as well as for two hypothetical imagers with reduced amplifier noise. The primary quantum efficiency for the detector is approximately 0.57; losses due to energy absorption noise and additive sources reduce the detective quantum efficiency to ~0.41 for exposures above 1 mR. For exposures below 0.1 mR, the detective quantum efficiency falls rapidly for amplifier noise values comparable to that found in the EG&G detector. Thus for thicker/denser objects [e.g., a pelvis (~30 cm water)] resulting in significantly reduced dose to the detector (e.g., ~0.001 mR)

improvements in amplifier noise (and/or x-ray converter, e.g. CsI;TI) will significantly improve detective quantum efficiency.

The temporal stability of the detector dark signal is presented in FIG. 5(c). This plot corresponds to a selected group of 'typical' pixels. The dark signal drifts significantly during the first 2 h of operation, which correlates with the change in temperature within the flat panel imager enclosure. After the temperature has stabilized, the dark signal also stabilizes. Based on these results, all cone beam computerized tomography scans were performed after the array had been powered-on for at least 2 hours. In some regions of the array, the dark signal does not stabilize, even after thermal equilibrium. It is assumed that these regions are the result of variations in the array manufacturing process.

The continuously changing scene in computerized tomography necessitates a detector with rapid read out and minimal temporal blurring, or 'lag.' Such characteristics have been measured using a short, intra-frame, x-ray exposure. FIG. 5(d) shows the pixel signal following a single radiographic exposure applied within the acquisition period of frame number 0. Subsequent frames exhibit lag signal ranging from ~4% to ~0.04% for frame members 1 through 9. It is interesting and important to note that the lag demonstrates a dependence not upon frame time, but also exclusively upon the number of frames.

Prior to reconstruction, the projections are corrected for stationary pixel-to-pixel variations in offset and gain. Defective pixels with significant variations in dark field signal or with aberrant signal response are median filtered. The resulting projections are padded by an additional 128 columns prior to reconstruction. The value of the padded pixels is set row-by-row to the average of the 7 pixels at the periphery of the array. Finally, to account for small variations in x-ray tube output, the signal in each projection is normalized using signal measured from the bare-beam monitors pixels mentioned above (nine pixels). The pre-construction processing can be performed on a 250 MHz UltraSparc processor, such as the Enterprise 450, Sun Microsystems, Sunnyvale, CA.

Feldkamp's filtered back-projection algorithm can be used to reconstruct the data set. Images are reconstructed on a Cartesian matrix of voxels $561 \times 561 \times N$, where the number of slices, N , depends on the object of interest. The voxel size used in these reconstructions was typically
5 0.25 x 0.25 x 0.25 mm. The filtering used in the reconstruction follows the formalism of Webb. Table 1 contains the three parameters that specify the filter used in these investigations. Upon completion of the reconstruction, an offset and scale parameters are constant for a 9 mm set of reconstruction and acquisition parameters. The reconstruction of the volumetric cone beam
10 computerized tomography data sets is also performed on the UltraSparc system.

The uniformity of response of the imaging system 300 over the three-dimensional (3-D) field-of-view (FOV) was studied by imaging a cylindrical water bath [110 mm diameter]. Scans of the same phantom were also
15 acquired on the conventional scanner. The response was examined along both radial and vertical profiles through the reconstructed volume.

The noise in reconstructed images of the water bath was studied as a function of x-ray exposure. Images were acquired at exposures of 131, 261, 653, 1310, 3260, and 6530 mR. The images were reconstructed on a
20 $561 \times 561 \times 11$ matrix with voxel dimensions of 0.25 mm on a side. For all reconstructions, the reconstruction filter was fixed at the parameters specified in Table 1. Varying these parameters can have a significant effect on the noise characteristics of the reconstructed images. The noise characteristics of these image sets were analyzed by analysis of the standard deviation in CT
25 number in $5 \times 5 \times 1$ regions throughout the data set, and by calculation of the noise power spectrum (NPS) from the 3D data sets. Both methods of analysis were performed as a function of exposure. The relative stability of the noise was assessed by examining the uniformity of the noise over the entire 3-D data set. These results indicated that the noise characteristics of
30 the data set vary only slightly with location. These initial results lend support to the application of noise power analysis, since stability is a necessary condition for proper interpretation of noise power results.

The noise-power spectrum (NPS) was analyzed from the volumetric data by extension of methods employed for analysis of known 2-D projection images. The volume data was normalized such that the mean CT number within the water cylinder was 1000. A tetragonal region (256 x 256 x 20 voxels) within the water cylinder was cropped from the volume, and a small number of voxel defects (always <1%) were 3 x 3 median filtered. In order to obtain a convergent 2-D central slice of the 3-D Fourier transform, the twenty slices were averaged along the z-direction, and it was found that averaging more slices did not affect the noise-power spectrum, i.e., the data was convergent. A background slice formed from the average of 81 slices in a separate scan was subtracted in order to reduce background trends. Low-frequency trends were further reduced by subtraction of a planar fit to the data, yielding a 2-D zero-mean realization. The two-dimensional Fast Fourier Transform (FFT) was computed from ensembles of sixteen 64 x 64 non-overlapping regions within the realization, and the results were averaged. The results were normalized to account for voxel size and for average in z, and the volume under the noise-power spectrum was compared to the square of the standard deviation. The resulting noise-power spectrum represents a central slice in the (u_x, u_y) domain, i.e., the Fourier counterpart to the (x, y) domain. Strips along the u_x axis were extracted in order to show 1-D power spectra, $NPS(u_x)$, e.g., are various exposure levels.

The noise characteristics of the cone beam computerized tomography system 300 were compared to those of the conventional computerized tomography scanner. To allow meaningful comparison, the two systems must demonstrate identical response over the range of signal variation. The response was tested by scanning an electron density phantom (shown in FIG. 6(b)) with the two systems. Seven inserts with coefficients near that of water were inserted into a 110 mm diameter water bath. The inserts are taken from the RMI electron density phantom having nominal CT numbers. In FIG. 6(b), clockwise from the top: CT Solid Water (CT#1001), BR-SRI Breast (CT#945), BRN-SR2 Brain (CT#1005), C133 Resin Mix (CT#1002), LV1 Liver (CT#1082), and, Polyethylene (CT#897). This phantom was imaged at

equivalent exposure and kVp with both the cone beam computerized tomography system 300 and the conventional scanner.

5 The attenuation coefficients (relative to water) reported by the cone beam computerized tomography system 300 were compared to those reported by the conventional scanner. A first-order fit to the measured data was calculated to determine the relative linearity of the two systems. The noise characteristics of the conventional scanner were also measured using the water cylinder test phantom described above images were acquired at 100kVp with a slice thickness of 1 mm at four different exposure levels (743,
10 1490, 2970, and 5940 mR). Three images were acquired at each exposure level. Reconstructions were performed on the conventional scanner using the 'High Res Head (#1 H)', 'Standard Head (#2)', and 'Smooth Abdomen (#3)' filters. The noise analysis was identical to that applied to the cone beam computerized tomography data sets. In order to compare noise results
15 measured on each system, analysis of the cone beam computerized tomography data sets was repeated wherein the cone beam computerized tomography data was first average over $2 \times 2 \times 4$ voxels to yield an equivalent ($0.5 \times 0.5 \times 1$ mm') voxel size to that given by the conventional scanner.

20 The spatial frequency transfer characteristics of the cone beam computerized tomography system 300 were measured using a wire test object, shown in FIG. 6(c). The test object consists of a 0.254 mm diameter steel wire suspended in a 50 mm diameter water bath. The phantom was imaged on the cone beam computerized tomography system 300 (at 100 kVp) with the wire centered on the axis of rotation 320 and with the wire located -30
25 mm off-axis. The resulting images were reconstructed on a high resolution reconstruction grid of $0.1 \times 0.1 \times 0.25$ mm³ using the filter described in Table 1. Six adjacent slices (each 0.25 mm thick) were averaged to generate a low noise point spread function (PSF). Orthogonal slices through the 2-D modulation transfer function (MTF) were calculated by first computing the
30 Radon transform of the point spread function (i.e., integrating along either the x or y axis), and then calculating the 1-D Fourier transform. Each 1-D profile was normalized to unity area. A correction was applied to compensate for the

finite diameter of the steel wire. For purposes of comparison, the same tests were performed on the conventional scanner at 100 kVp for a slice thickness of 1.5 mm. Images were reconstructed using three different reconstruction filters ["High Res Head (#1 H)," "Standard Head (#2)," and "Smooth Abdomen (#3)"].

The relative imaging performance of the cone beam computerized tomography system 300 and the conventional scanner were compared using phantoms and small animals. A simple comparison in soft-tissue detectability was performed with the phantom shown in FIG. 6(b). The proximity in CT number between each of the six cylinders makes this phantom a useful test object for examining contrast sensitivity and soft-tissue detectability, images were acquired of the phantom with both the cone beam computerized tomography system 300 and conventional scanners. Multiple high-resolution

cone beam computerized tomography slices were averaged to produce an equivalent slice thickness to that used on the conventional scanner (1.5 mm). Equivalent exposure (2980 mR) and kVp were used in the two different scans.

A second test of soft-tissue sensitivity was performed by imaging a laboratory rat that had been euthanized for other purposes, FIG. 6(d). A scanning procedure identical to that described above was used, delivering an in-air, on-axis exposure of 2980 mR at 100 kVp for both systems. The resulting 3-D data was reconstructed at voxel sizes of $0.25 \times 0.25 \times 0.25 \text{ mm}^3$. The subject was also scanned on the conventional computerized tomography scanner at a slice thickness of 1.5 mm. This scan delivered the same imaging dose as was delivered by the cone beam computerized tomography system 300. For purposes of intercomparison, six slices from the cone beam computerized tomography data set were averaged to produce a slice thickness equivalent to that of the conventional scan. The imagers were displayed at comparable window and level to allow comparison.

The uniformity of response of the cone beam computerized tomography scanner shown in shown in FIGS. 7(a) - 7(d). Axial and sagittal slices through the cone beam computerized tomography 3-D data set are

shown. The images demonstrate a relatively uniform response over the entire field of view of the system. A slight non-uniformity of approximately 20 CT numbers (2%) is visible in the histogram equalized- regions of the images. This non-uniformity appears as a combined cupping and capping artifact. The radial profile (FIG. 7(c)) illustrates this point further by comparing to the results obtained from the conventional scanner (dotted line). An internal check of the reconstruction process using simulated projection data demonstrates that the non-uniformity is an artifact of the reconstruction process and is dependent upon the choice of filtering parameters. Apart from the non-uniformity inherent to the reconstruction, the response of the cone beam computerized tomography system 300 is highly uniform, particularly along the z-dimension.

In addition to demonstrating uniformity of system response, the images in FIG. 7 also demonstrate uniform noise characteristics with few artifacts. This is the case for the full range of exposures studied. The magnitude and uniformity of the noise is demonstrated in FIGS. 8(a) - 8(d). The noise varies to a slight degree along the radial axis and to a negligible degree along the vertical axis. A slight dependence on radial position is expected due to the differences in transmission across the cylindrical water bath. FIG. 8(c) also presents the measured dependence of noise on exposure [also shown below, in relation to FIG. 9(b)]. Overall, the cone beam computerized tomography system 300 is capable of achieving a noise level of approximately 20 CT numbers for an in-air exposure of 6560 MR at isocenter.

The noise measured for the cone beam computerized tomography system 300 as a function of exposure is shown in the top curve of FIG. 9(b). The noise is seen to decrease from -80 units at the lowest exposure examined down to -20 units at the highest. Superimposed is a least squares fit of the form $\sigma = a + b / \sqrt{X}$, where σ is the noise in voxel values, X is the exposure in air at the isocenter, and a and b are constants obtained from the numerical fit. This inverse-square root dependence upon exposure is consistent with basic noise transfer theory for x-ray tomographic reconstructions.

In order to examine the linearity and accuracy of system response, the CT numbers reported by the cone beam computerized tomography system 300 for a variety of materials (FIG. 6) were compared to those reported by the conventional scanner. As shown in FIG. 9(b), the CT numbers of the cone beam computerized tomography system 300 agree well with those of the conventional scanner. The largest discrepancy over the range of CT numbers was 8 units, with an average discrepancy of 5.7. The high coefficient of correlation indicates that, over the range examined, the values reported by the cone beam computerized tomography system 300 are proportional to attenuation coefficient.

The voxel noise of the cone beam computerized tomography system 300 and the conventional scanner was compared as a function of exposure, shown in FIG. 9(b). Shown by the open circles and dashed lines are the results for the conventional scanner using the "High-Res Head (#1H)" and "Standard Head (#2)" reconstruction filters. In each case, the noise decreases with exposure. An exact comparison between the two systems requires that both data sets be reconstructed at equivalent voxel size and with the same reconstruction filter. The requirement for equivalent voxel size was achieved by repeating the noise analysis for the cone beam computerized tomography system 300, with the volume data averaged to give a voxel size equivalent to that of the scanner.

In order to illustrate the effect of the reconstruction filter upon the voxel noise, reconstructions were performed with both the "High-Res Head" and "Standard Head" reconstruction filters. The noise for the cone beam computerized tomography system 300 at equivalent voxel size is shown by the lower solid curve with a least-squares fit superimposed. At equivalent voxel size, it is clear that the cone beam computerized tomography system 300 has higher noise at lower exposures than the "Standard Head" computerized tomography scanner results. Compared to the "High-Res Head" results for the conventional scanner, however, the cone beam computerized tomography system 300 actually provides lower noise at all but the very highest exposures. Clearly, careful matching of reconstruction filters

and reconstruction matrix is required to permit exact intercomparison of the two systems. Nonetheless, the results obtained using the cone beam computerized tomography system 300 are encouraging, since the early prototype flat-panel detector used in this system is known to exhibit a fairly high level of additive electronics noise, a factor of -5-10 higher than that achieved, by more recent electronics designs.

Results of the noise-power spectrum measurements are summarized in FIGS. 10(a) - 10(c). The 2-D noise-power spectrum in the axial plane (FIG. 10(a)) exhibits a spectral shape typical of systems employing filtered back-projection reconstruction. The spectral density is reduced (but non-zero) near zero-frequency, increases at mid-frequencies due to the ramp filter (e.g., peaking around -0.5 mm^{-1}), and declines at higher frequencies by virtue of the low-pass noise characteristics of the system (e.g., 2-D image blur and choice of apodisation window). Slices of the noise-power spectrum along the u_x dimension are shown in FIG. 10(b) for various exposure levels. Since the mean signal level is fixed for each case (i.e., CT# = 1000 within the water phantom), the noise-power spectrum decreases with increasing exposure. Specifically, the noise-power spectrum appears inversely proportional to exposure in a fashion consistent with the form of the numerical fits in FIG. 9(b). As shown in FIG. 10(c), the noise-power spectrum measured at -1.3 R (in air at isocenter) is -30 mm^3 near zero-frequency, increases by a factor of -4 at mid-frequencies, and then descends to about the initial level of spectral density at the Nyquist frequency.

Superimposed in FIG. 10(c) are the results measured for the conventional scanner using three reconstruction filters, and to facilitate intercomparison, noise-power spectrum results for the cone beam computerized tomography system 300 are shown for an equivalent voxel size. For the #2 and #3 filters, the conventional scanner exhibits a noise-power spectrum with the characteristic shape described above; however, the high-resolution #1 H filter is seen to significantly amplify high-frequency noise. The cone beam computerized tomography system 300 appears to exhibit low-frequency noise-power spectrum comparable to the conventional scanner

using the #2 and #1 H filters. Given that the choice of reconstruction filter can significantly affect noise and spatial resolution, and considering the two cases that seem most closely matched the cone beam computerized tomography system 300 - even in its initial, un-optimized configuration -- appears to provide noise performance comparable to the conventional scanner. As evident in FIG. 9(b), the cone beam computerized tomography system 300 exhibits lower voxel noise than the conventional scanner (#1H) at low exposures. Similarly, the cone beam computerized tomography system 300 exhibits reduced high-frequency noise-power spectrum. These initial results are especially promising considering the on-going improvements in FPI design and readout electronics.

The response of the cone beam computerized tomography system 300 to the wire test object is presented in FIG. 11(a). Overall, the PSF is symmetric (aside from a small streak artifact believed associated with the image lag characteristics of the system) and has a full-width at half-maximum (FWHM) of 0.6 mm. The system MTF is shown in FIG. 11(b) for both the on- and off-axis wire results. These results suggest that the frequency pass of the system in the $z = 0$ plane does not change significantly over the relatively small (-30 mm) range examined. The strong influence of the reconstruction filter is demonstrated in the MTF results for the conventional scanner, also shown in FIG. 11(b).

The "Standard Head (#2)" filter significantly reduces the signal pass of the system compared to the High-Res Head (#1 H) filter. The results demonstrate that the MTF of the conventional scanner is comparable to that of the cone beam computerized tomography system 300 when the "High-Res Head (#1 H)" filter is used. This observation is consistent with the noise results presented in FIG. 9(b). The resolution of the cone beam computerized tomography system 300 and conventional scanner have not been compared in the z -dimension. It is expected, however, that the spatial resolution of the cone beam computerized tomography system 300 in the z -dimension will be comparable to that measured in the axial plane. Of course, the spatial resolution of the conventional scanner will be limited by the selected slice

thickness, which is typically 1 mm or greater. The nearly isotropic resolution of the cone beam computerized tomography system 300 is expected to be a significant advantage for detection and localization.

FIGS. 12(a) and 12(b) show axial image slices of the low-contrast phantom obtained on the cone beam computerized tomography system 300 and the conventional computerized scanner at equivalent kVp and exposure. The grayscale window in each case is quite narrow in order to maximize the displayed contrast, and despite the slight signal non-uniformity evident for the cone beam computerized tomography image (cupping/capping artifact discussed above) the visibility of each insert is comparable to the conventional scanner. The mean signal values for each material are as shown in FIG. 9(a). Slight differences in system response (e.g., due to detector response, x-ray spectrum, etc.) can result in contrast reversal for materials with CT# very close to that of water. For example in the case of the brain insert (lower right), even the slight (-5 CT#) difference between the mean value reported by the cone beam computerized tomography system 300 and the conventional scanner is sufficient to give an apparent inversion in the density of the material relative to water. The minimum detectable contrast is arguably superior for the cone beam computerized tomography system 300 (e.g., visibility of the brain and CB-3 inserts), but this remains to be verified by a more controlled, quantitative observer study.

The overall performance of the cone beam computerized tomography system 300 is demonstrated in the images of the volumetric data set illustrated in FIGS. 13(a)-13(i). These images of an euthanized rat demonstrate the soft tissue sensitivity and high spatial resolution of the system. Example images are shown from various regions throughout the volumetric set [e.g., in regions of the lungs (a,b,c), the kidney (d,e,f), and lower spine (g,h,i)] to illustrate the quantity and uniform quality of the data produced with the cone beam computerized tomography system 300. The clear visualization of soft-tissue structures demonstrates the soft-tissue contrast sensitivity of the scanner.

In FIGS. 13(a) -13(c), the window and level have been set to emphasize features in the lung of the rat. In addition to the lung detail, there are some streak artifacts evident, the origin of which is unknown, but is believed to be associated with detector lag effects or beam hardening.

5 The soft tissue contrast sensitivity of the cone beam computerized tomography system 300 is illustrated in FIGS. 13(d) -13(f), in which the window and level have been set to delineate fat and muscle. The cross-hair in each image indicates the location of the rat's left kidney. These images illustrate the advantage of a nearly isotropic spatial resolution for delineation
10 of a 3-D structure such as the kidney. Other structures, such as the stomach, bowel and liver are also clearly visible.

 The spatial resolution performance of the system 300 is demonstrated in FIGS. 13(g-i), in which the same rat data set is displayed with window and level selected to display bony features. The clear visibility of the intervertebral
15 spaces and the non-cortical bone in the pelvis is stunning. It should be kept in mind that this level of detail was produced on a cone beam computerized tomography system 300 that operates on a scale that mimics the geometry of the linear accelerator. Therefore, this level of detail would be expected in the clinical implementation of the device, given accurate correction of mechanical
20 flex. The volumetric data set is illustrated further in FIG. 14, in which volume renderings demonstrate the fully 3-D nature of the data set and show the level of detail contained within the cone beam computerized tomography data. It is interesting to note that all the data presented in FIGS. 13 and 14 were
 obtained from a single acquisition performed in a single rotation.

25 Finally, the quality of images produced by the cone beam computerized tomography system 300 was assessed by comparison to images produced by the conventional scanner. FIGS. 15(a) - 15(b) show an axial slice of the rat acquired on the two systems. At equivalent exposure, the images produced by the two systems are of comparable quality both in terms of spatial
30 resolution and contrast sensitivity. The flat panel imager-based cone beam computerized tomography image exhibits exquisite spatial resolution and provides clear delineation of soft-tissue boundaries and detail in the gut. The

spatial resolution of the cone beam computerized tomography system 300 appears to exceed that of the conventional scanner; however, it must be noted that restrictions in available reconstruction matrices for the conventional computerized tomography scanner limited the voxel size to twice that of the cone beam computerized tomography image. Lack of obvious pixelation in the flat panel imager-based cone beam computerized tomography image indicates that this level of detail represents the physical limits in spatial resolution of the current system.

The objective of these investigations is to evaluate the applicability of flat-panel technology as a detector in a cone beam computerized tomography system, specifically, a tomographic imaging system for use in the guidance of radiation therapy on a medical linear accelerator.

The quantitative and qualitative results of our studies suggest that a cone beam computerized tomography scanner based on flat panel detector technology is a viable means for high performance computed tomography. Initial studies of signal response uniformity demonstrated that the response of the system is uniform over the field of view to within $\pm 2\%$, with the slight degree of non-uniformity apparent as a combined cupping and capping artifact in the x-y plane attributable to a reconstruction artifact. The linearity of response was demonstrated using a range of soft-tissue test materials and was found to be linear to within $\pm 0.6\%$. Measurements of image noise versus exposure demonstrate that the prototype cone beam computerized tomography system 300 performs comparably to the conventional scanner, demonstrating the inverse square root exposure dependence predicted by theory. Investigations of noise power spectrum and spatial frequency response for the two systems reinforce these conclusions and illustrate the advantages of developing more extensive (empirical and theoretical) frequency-dependent characterization methods for volumetric computed tomography systems.

In addition to the quantitative measures of performance, the images of low-contrast phantoms and small animal anatomy confirm the conclusions

drawn from these measures, showing excellent detail and soft-tissue contrast, more than sufficient for tissue localization in radiation oncology.

The results presented here demonstrate the potential of this approach for volumetric imaging. However, this study has been performed under conditions of small object size and small cone angle. These conditions are imposed by the size of the detector used in this investigation. Imaging with larger detectors allows increased cone angle and, for computerized tomography, increased object thickness. The extrapolation of performance based on the results presented here to that for larger detectors must be done with some caution. Imaging larger objects with an increased field of view will result in increased scatter and reduced transmission. The increase in scatter can be expected to have a negative impact on computerized tomography imaging performance by introducing non-uniformities in the reconstructed image (e.g., cupping and/or streaks), and by adding additional x-ray quantum noise to the image signal. The magnitude of scatter reaching the detector will depend greatly on the cone-angle and air gap employed, and studies suggest that scatter at these distances may be reduced compared to conventional radiographic applications. Quantifying the magnitude of the x-ray scatter problem and developing methods to reduce it are areas of ongoing investigation.

In addition to concerns of x-ray scatter at large cone-angles, the scanning of larger objects will significantly reduce the fluence arriving at the detector. This reduced transmission will negatively impact the performance of the flat-panel detector. Currently available flat panel imagers demonstrate performance inferior to conventional image intensifiers at fluoroscopic exposure rates, due to the presence of additive noise in the flat-panel readout electronics. Additive noise causes the detected quantum efficiency of the imager to depend on the number of x-rays forming an image. This dependence is illustrated in FIG. 16 for the flat-panel imager 306 used in these investigations and for hypothetical detectors that embody the most recent advances in imager 306 design, including higher x-ray quantum

detection efficiency through the use of CsI:TI and a reduction in additive noise through improvements in readout electronics.

5 The zero-frequency detected quantum efficiency was computed using a model for signal and noise transfer that has demonstrated excellent agreement with measurements. It is clear from FIG. 16 that improvements in the x-ray converter and electronics noise significantly reduce the exposure dependency of the detected quantum efficiency over the broad range of exposures required for computerized tomography. The magnitude of the reduction depends greatly on the amplifier noise in the system. For the 10 prototype imager used in these studies, the amplifier noise is very high at 12,700 e. For the low transmitted exposure levels in computerized tomography of pelvic anatomy, for example, this detector would achieve a zero-frequency detected quantum efficiency of less than 10%. In comparison, an imager that incorporates the recent advances in design listed above (e.g., 15 a high-quality CsI:TI converter and amplifier noise of 3000 e or better) would achieve a higher detected quantum efficiency (~65%) at full transmission and maintain a detected quantum efficiency of >40% even at the low exposure levels. Such enhancements in imager design are within the current capabilities of flat panel imager manufacturers and will greatly facilitate the application of flat 20 panel imagers in cone-beam computerized tomography of human beings. Furthermore, these improvements are largely driven by other forces in digital imaging that anticipates use of flat panel imagers in place of conventional image-intensifier systems for interventional fluoroscopy. For this reason, it can be expected that imagers with such characteristics will be available within the 25 next five years.

Overall, the operating characteristics of the flat-panel are highly compatible with acquisition in a cone beam computerized tomography scanning geometry. Unlike image-intensifier or lens based systems, flat panel detectors are geometrically robust under a rotating geometry, eliminating concerns of 30 image distortion. The proximity of the analog-to-digital converter to the pixel element and the relatively large charge signals make the panels robust in high radio-frequency power environments; this is of particular interest for radiotherapy

applications. The high readout rate of these detectors allows for imaging sequences of 300 projection images to be acquired within 10 seconds (operating at 30 fps). This is more than sufficient to satisfy the allowable rotation rates for the gantry of a medical linear accelerator. In fact, while the International Electromechanical Commission (IEC) recommends less than 1 revolution per minute for linear accelerators, it would be reasonable to reconsider such constraints in light of the advantages of cone beam computerized tomography guidance in the treatment room. Currently, the detector size and aspect ratio are driven by the needs of digital radiography, producing detectors comparable in size to radiographic film. These sizes limit the field-of-view of the reconstruction if sufficient clearance is to be maintained between the detector and patient during gantry rotation. This problem can be addressed using offset detector schemes that use 360° of gantry rotation. Ultimately, a specialized detector could be designed with a size and aspect ratio that match the requirements for cone beam computerized tomography (e.g., a $-25 \times 50 \text{ cm}^2$ area panel).

Given the potential that this technology is demonstrating, the opportunities for new areas of application for computed tomography are significant. Imaging systems based on this technology can be constructed to address specific imaging problems, including non-destructive testing (at kilovoltage or megavoltage energies), early detection and monitoring of specific medical conditions, and, of course, navigational imaging for therapies. The compact nature of the panels allow flat panel imager-based cone beam computerized tomography imagers to be applied in situations that would never be considered feasible for a conventional computerized tomography scanner. The cone beam computerized tomography approach offers two important features that dramatically reduce its cost in comparison to a conventional scanner. First, the cone-beam nature of the acquisition does not require an additional mechanism to move the patient (or object) during image acquisition. Second, the use of a cone-beam, as opposed to a fan-beam, significantly increases the x-ray utilization, lowering the x-ray tube heat capacity required for volumetric scanning. For the same source and detector geometry, the efficiency roughly scales with the slice thickness. For example, the x-ray utilization increased by a factor of 30 in going from a 3 mm slice in a conventional scanner to a cone-angle

corresponding to a 100 mm slice with a cone-beam system, This would decrease heat-load capacities dramatically. From our experience, a 5200 kHU x-ray tube costs approximately \$70,000, whereas a 600 kHU x-ray tube (a factor of -10 lower in capacity) costs roughly \$6,000.

5 Cone-beam computed tomography has been a topic of active research and development for over a decade in areas such as nuclear medicine and industrial testing; however, only recently has it begun to appear in the diagnostic computerized tomography arena. The developments in this area have been for the most part limited to multi-slice detectors. In this investigation, the use of an
10 alternative detector for high-quality computerized tomography has been studied. The results of the investigation suggest that there is a significant potential for the use of these detectors in cone beam computerized tomography systems for radiotherapy and quite possibly for diagnostic and interventional computerized tomography imaging tasks that will take advantage of the fully 3-D nature of
15 cone beam computerized tomography.

 Based upon the positive results presented previously with respect to the cone beam computerized tomography system 300, several embodiments of a flat panel imager-based kilovoltage cone beam computerized tomography scanner for guiding radiation therapy on a medical linear accelerator are
20 envisioned. For example, FIGS. 17(a)-(e) and 18 are diagrammatic and schematic views of an embodiment of a wall-mounted cone beam computerized tomography system 400. The cone beam computerized tomography system 400 includes an x-ray source, such as x-ray tube 402, and a flat-panel imager 404 mounted on a gantry 406. The x-ray tube 402
25 generates a beam of x-rays 407 in the form of a cone or pyramid that have an energy ranging from approximately 30KeV to 150 KeV, preferably approximately 100 KeV. The flat-panel imager 404 employs amorphous silicon detectors.

 The system 400 may be retrofitted onto an existing or new radiation
30 therapy system 700 that includes a separate radiation therapy x-ray source, such as a linear source 409, that operates at a power level higher than that of x-ray tube 402 so as to allow for treatment of a target volume in a patient.

The linear source 409 generates a beam of x-rays or particles 411, such as photons or electrons, that have an energy ranging from 4 MeV to 25 MeV. The system 400 may also include an imager (not shown) that is aligned with the linear source 409 with the patient interposed therebetween. The imager forms projection images of the patient based on the remnants of the beam 411 that passes through the patient. Note that the x-ray sources 402 and 409 may be separate and contained with the same structure or be combined into a single source that can generate x-rays of different energies.

As shown in FIGS. 17(a)-(e) and 18-19, the flat-panel imager 404 can be mounted to the face of a flat, circular, rotatable drum 408 of the gantry 406 of a medical linear accelerator 409, where the x-ray beam 407 produced by the x-ray tube 402 is approximately orthogonal to the treatment beam 411 produced by the radiation therapy source 409. Attachment of the flat plane imager 404 is accomplished by an imager support system 413 that includes three 1m long arms 410, 412 and 415 that form a tripod. Side arms 410 and 415 are identical to one another in shape and have ends attached to a Ax95 Guy pivot 417 which in turn is attached to a mounting 414 by screws that are threaded through aligned threaded holes of the pivot 417 and threaded holes 425 and 431 of plates 433 and 435, respectively, as shown in FIGS. 18 and 19(a)-(b). As shown in FIGS. 17(b) and 18, the mountings 414 for the arms 410 and 415 are aligned with one another along a line segment 419 that is contained within a plane 421 that is parallel to and offset by approximately 30 cm from the plane containing the flat-plane imager 404. The mountings 414 are separated from one another by approximately 70 cm and are symmetrically positioned with respect to a plane bisecting an imager mount 423 that is attached to the drum 408 270° from the radiation therapy source 409.

As shown in FIGS. 18 and 19(a)-(b), each mounting 414 is attached to an end portion 416 of the drum 408 by inserting a threaded male member 418 through an opening 437 formed through the drum 408. Once inserted, the male member 418 is attached to the drum 408 by tightening a nut 420 onto the threaded male member 418. The other ends of the arms 410 and 415 are attached to Ax95 Guy pivots 422 attached to the back of an 3/8 inch thick

Aluminum square plate 424 is attached to the rear of the flat-panel imager 404 via bolts (not shown).

As shown in FIGS. 17(d)-(e), there are two preset positions of the flat panel imager 404 relative to the plate 424. As shown in FIG. 17(d), the flat panel imager 404 is centered about the ends of the arm 412. In order to provide a larger field of view, an offset flat panel imager 404 can be used as shown in FIG. 17(e) where the imager 404 is attached to a side of the plate 424 via bolts. Note that it is possible to use a motorized system to move the flat panel imager 404 relative to the plate 424 to provide an easy way to vary the field of view of a cone beam computerized tomography system.

A center arm 412 is also attached to the drum 408 and the flat-panel imager 404. The center arm 412 has one end attached to Ax95 Guy pivot 427 that is in turn attached to a tapped, triangular-shaped, reinforcing plate 426 formed on the drum 408 as shown in FIGS. 17(b) and 18. The plate 426 is approximately 433.8mm from the rotational axis 428 that intersects the iso-center 430 of the imaging system 400. A second end of the center arm 412 is attached to the plate 424 via a Cx95A right angle joint 425.

As shown in FIGS. 17(b) and 18, the end of the arm 412 lies along a line that is the perpendicular bisector of the line segment 419 and is radially separated from the midpoint between mountings 414 as measured along line segment 419 by a distance D of approximately 30 cm.

As shown in FIGS. 17(b) and 18, the other ends of the arms 410, 412 and 415 are attached to the plate 424 so as to be positioned approximately 20 cm from the rear edge 429 of the plate 434 and approximately midway between the left and right edges of the plate 434.

Once the arms 410, 412 and 415 are attached to the drum 408 and the plate 424, the arms can be pivoted so that the flat panel imager 404 moves to a position where its rear side is separated from the iso-center 430 by a distance L of approximately 600 mm. One advantage of the imager support system 413 is that it can be used to retrofit existing stand-alone radiation treatment devices so they have the capability to have a flat panel imager attached thereto. The imager support system 413 is very rigid, i.e., constant

tension and compression, which reduces movement of the imager 404 and so leads to cleaner imaging data.

Note that the x-ray tube 402 can also be retrofitted onto an existing stand-alone treatment device so as to be positioned opposite to the flat panel imager 404. As shown in FIGS. 17(a)-(e), the x-ray tube 402 is attached to tube support 440 that is composed of a pair of front and rear faces 442 and 444 and a pair of side faces 446. A multi-leaf collimator 448 is supported within the interior of the tube support 440. The front and rear faces 442 and 444 each include three openings 450, 452 that are aligned with one another and receive three cylindrical support arms 454 that are attached to a bearing housing 456 that is bolted to the drum 408. The tube support 440 and the x-ray tube 402 are able to slide along the support arms 454. Note that a cable support 458 spans between the tube support 440 and the bearing housing 456 and contains the wiring necessary to operate the x-ray tube 402.

An alternative imager support system for the flat panel imager 404 of FIGS. 17 is shown in FIGS. 20(a)-(b). In particular, the imager support system 507 shown in FIGS. 20(a)-(b) includes a single pivoting arm 510 that has one end 511 pivotably attached to a lower corner of the radiation therapy source 409. The other end 512 of the arm 510 is pivotably attached to an end of the flat-panel imager 404. The arm 510 and flat-panel imager 404 are movable from a retracted position of FIG. 20(a) to an extended position of FIG. 20(b) and vice versa. Movement of the arm 510 and the flat-panel imager 404 may be moved either manually or via a motor.

Note that when the imager support system 507 is used, the x-ray tube 402 is attached to a second lower corner of the radiation therapy source 409 in order to simplify the support structure and reduce the mechanical complexity of the overall system. The position of the x-ray tube 402 also reduces interference with staff access to the patient. Note that in this embodiment, the distance from the x-ray tube 402 to the axis of rotation 428 is not necessarily equal to the distance from the radiation therapy source 409 to the axis of rotation 428. Also, the amount of extension of the arm 510 shown in FIG. 20(b) will vary depending on the desired field of view for cone beam

computerized tomography imaging. Note that if the mechanics are engineered to be sufficiently precise, the arm 510 can move in and out during image acquisition during gantry rotation so as to allow the imager 404 to dynamically avoid potential rotation-induced collisions with the patient or the table. The head of the radiation therapy source 409 can be altered to provide additional lead shielding on the imager side to limit radiation induced damage to the imager 404 while in the retracted position of FIG. 20(a). This will increase the life span of the imager 404.

A second alternative imager support system for the flat panel imager 404 of FIG. 17 is shown in FIGS. 21(a)-(b). In particular, the imager support system 607 shown in FIGS. 21(a)-(b) includes a single C-arm 610 that is attached to an arm support 611 that is attached to the front or rear of the radiation therapy source 409. At one end of the C-arm 610 the x-ray tube 402 is attached and at the other end the flat-panel imager 404 is attached. The C-arm 610 is moved either manually or by a motor within the arm support 611, so that the x-ray tube 402 and the flat-panel imager 404 can move along an arc.

Note that in this embodiment, the distance from the x-ray tube 402 to the axis of rotation 428 is not necessarily equal to the distance from the radiation therapy source 409 to the axis of rotation 428. The arm 610 does not necessarily be in the shape of an arc of a circle. Also, the axis of rotation of the arm 610 is not necessarily coincident with the axis of rotation 428 of the radiation therapy source 409, which allows the same device to be fitted on machines with different face-to-isocenter distances without altering the radius of the C-arm 610.

Use of the C-arm 610 of FIGS. 21(a)-(b) allows for a great amount of flexibility in obtaining cone beam computerized tomography images. For instance, image data can be obtained by only having the drum 408 of the gantry 406 rotate. Image data can be obtained in a second manner by having the C-arm 610 move independently of the gantry 406 in a circular path. Image data can be obtained by having the C-arm 610 and the drum 408 work cooperatively to generate images along a circular path so that the angular

range of acquisition is increased and so instabilities in the angular velocity of the gantry are addressed. A fourth manner of imaging involves rotating the drum 408 and pivoting the C-arm 610 about the mounting point on the gantry 406 with a sinusoidal pattern to effect non-circular orbits that involve a sinusoidal trajectory on a spherical surface. Such a non-circular orbit allows more complete image reconstructions by satisfying Tuy's condition.

FIG. 22 shows a portable cone beam computerized tomography system 700. In this embodiment, the system 700 is on a mobile platform 702 so that it can be moved relative to a patient 441 positioned on a table 443 relative to a rotating radiation therapy source 409 (not shown). The cone beam computerized tomography system 700 includes an x-ray source, such as x-ray tube 402 positioned on one side of a C-arm 704, and a flat-panel imager 404 positioned on an opposite side of the C-arm 704. The C-arm 704 can rotate about two axes of rotation when in operation. The system 700 can be moved to a radiation therapy system (not shown) and can be used to generate images that aid in the alignment of the radiation therapy system.

With the above descriptions of the cone beam computerized tomography system 400 and the various embodiments of the imager support systems shown in FIGS. 18-22 in mind, operation of the system 400 is described below. In the description to follow, the term "shape" of the radiation therapy beam 411 is understood to refer to the spatial distribution of the beam in a plane perpendicular to the direction of the beam or to the frequency modulation of the beam after being transmitted through some beam-limiting device. The term "planning image" refers to an image of the patient acquired by the cone beam computerized tomography system 400 prior to treatment delivery used for radiation therapy treatment planning. The term "constrained plan set" refers to a plurality of radiation therapy treatment plans for a given patient, where each radiation therapy treatment plan is calculated assuming some perturbation of lesion location and/or orientation compared to that in the planning image. For example, a constrained plan set could be calculated where each plan corresponds to a different magnitude of lesion rotation about the y and/or z axes.

The cone beam computerized tomography imaging system 400 preferably includes an x-ray tube 402 and a flat panel imager 404 in any one of the geometries illustrated in FIGS. 23(a)-(d), capable of forming 3-D images of the patient on the treatment table in the treatment position. The x-ray tube 402 may be operated so as to produce a pulsed or continuous beam of x-rays 407. The flat panel imager 404 includes an active matrix of imaging pixels incorporating mechanisms for: 1.) converting incident x-rays to electronic charge (e.g., a scintillator in combination with optically sensitive elements at each pixel, or a photoconductor); 2.) integrating and storing the electronic charge at each pixel (e.g., the capacitance of photodiode(s), capacitors, etc. located at each pixel); and 3.) reading the electronic charge out of the device (e.g., a thin-film transistor switch or the like at each pixel, with associated switching control lines and readout lines). The x-ray tube 402 and the flat panel imager 404 preferably move in a circular orbit (or variation thereof) about the longitudinal axis of the patient. Depending on which ones of the imager support systems used in FIGS. 18-22, the imager support system should accommodate offsets in the x and/or z directions as illustrated in FIG. 23(b). Note that the combined motion of the x-ray tube 402 and/or the flat panel imager 404 in x, y, and/or z is termed the orbit, and may be circular about the patient, or non-circular, e.g., comprising of some combination of linear, sinusoidal, circular, and/or random paths. For example, in the case where the source 402 and imager 404 move independently with respect to one another, the source 402 can move on a sinusoidal or sawtooth path constrained to the surface of a cylinder while the imager 404 moves in a circular path on the surface of a cylinder. In this scenario, the collimator adjusts in real time the shape of the radiation field so it is confined to the imager 404 despite the allowed independent motion of the source 402 and imager 404.

Cone beam computerized tomography image acquisition involves acquisition of a plurality of 2-D images, where each image preferably corresponds to a different orientation of the x-ray beam 407 and the flat panel imager 404 with respect to the patient 441, e.g., where the x-ray tube 402 and the flat panel imager 404 traverse a circular or non-circular path about the patient

441 as illustrated in FIG. 23(d). Note that the cone beam computerized tomography image is preferably acquired with the patient on the treatment table, in the treatment position, and immediately prior to treatment delivery. The processes involved in the preferred method for cone beam computerized tomography image acquisition are illustrated in FIG. 24, divided conceptually into a variety of off-line and on-line processes, and mechanisms for 2-D image acquisition and 3-D image reconstruction.

The off-line processes schematically shown in FIG. 24 include acquisition of a plurality of 2-D images acquired in the absence of x-ray irradiation (termed dark fields) and with uniform x-ray irradiation (termed flood fields). Such dark and flood fields are used to correct stationary nonuniformities in the imaging system arising from nonuniformity in pixel operational and response characteristics. Also included is a mechanism for identifying and correcting defective pixels in the 2-D images (e.g., a pixel defect map that identifies aberrant pixel coordinates, and application of a filter to the corresponding pixel values). Thirdly, a measure and process for correction of orbit non-idealities, described below, is preferably employed.

The on-line processes schematically shown in FIG. 24 include: 1.) control and monitoring of the x-ray tube; 2.) control and monitoring of the orbit traversed by the x-ray tube 402 and the flat panel imager 404 (e.g., by rotating the gantry 406); and 3.) control and readout of the flat panel imager 404. The x-ray source 402 produces x-rays in either a pulsed or continuous manner, and variations in the magnitude of x-ray tube output are monitored by an output monitor, which preferably includes a radiation sensitive electronic device such as a diode placed inside the x-ray tube collimator assembly. Alternatively, the output monitor could be placed outside the x-ray tube 402 in a position that allows it to measure variations in x-ray tube output, or the output could be measured using pixels on the flat panel imager 404, such that these pixels are not occluded by the patient in the plurality of 2-D projection images. The orbit of the x-ray tube 402 and the flat panel imager 404 about the patient is preferably controlled via computer-controlled rotation of the gantry 406, combined with a precise measurement of the gantry angle at which each 2-D image is acquired. For embodiments in

which the x-ray source 402 and the flat panel imager 404 are not both mounted on the treatment gantry 406, such as the portable embodiment of FIG. 22, a similar mechanism for measuring and recording the location of these two components for each 2-D image is employed. Thirdly, a plurality of 2-D images are read from the flat panel imager 404 by a control/acquisition computer. The readout of the flat panel imager 404 is preferably synchronized with the operation of the x-ray tube 402 as well as with the rotation of the x-ray tube 402 and the flat panel imager 404 support structure(s), such as those described previously with respect to FIGS. 18-22. The timing of x-ray exposures, gantry rotation, and flat panel imager readout is preferably synchronized by: 1.) the control/acquisition computer; or 2.) an external trigger mechanism (gating source), such as a device for active breathing control, electrocardiac gating, etc. For the former case, the preferred embodiment includes computer-control of: 1.) x-ray pulses generated by the x-ray source 402; 2.) gantry rotation (e.g., in increments of $\sim 1^\circ$ through $\sim 360^\circ$); and flat panel imager readout (e.g., at a readout rate consistent with the limitations in x-ray tube output and gantry rotation). For the latter case, the preferred embodiment is such that the gating source triggers x-ray production, gantry rotation, and flat panel imager readout in a manner synchronized with the motion of anatomical structures in the patient 441 in order to reduce the deleterious effects of organ motion in image reconstructions.

The preferred embodiment includes a mechanism (reconstruction engine) for high-speed cone beam computerized tomography image reconstruction. The plurality of 2-D projections is first processed by dark and flood field correction, and the measurements of orbit non-ideality (below), tube output variations, and gantry rotation are used together with the processed 2-D projections to form 3-D cone beam computerized tomography image reconstructions of the patient 441. A variety of cone-beam reconstruction techniques are known within the art, including cone-beam filtered back-projection. The cone beam computerized tomography image is then made available to a system for on-line treatment planning.

In the interim between the 2-D image acquisition and correction of lesion localization errors, the patient 441 is preferably monitored by periodic

radiographs obtained with the flat panel imager at one or more gantry angles. In the preferred embodiment, these monitor radiographs are analyzed (e.g., by calculation of difference images) in order to provide a check against intrafraction motion of the patient 441.

5 The preferred embodiment includes a computer-controlled treatment table 443 for correction of lesion localization errors. The table 443 preferably allows translation of the patient 441 in the x, y, and z directions as well as rotation about the x axis. Rotation about the y axis (tilt) and z axis (roll) is possible for an embodiment in which lesion localization errors are corrected by such motions (as
10 opposed to correction of such errors through selection of an appropriate RTTP from a constrained plan set), provided that such motions do not cause uncertainty in the location/orientation of the lesion 444 and/or surrounding structures, e.g., due to the effects of gravity. Furthermore, the treatment table 443 is preferably constructed of radio-translucent material so as not to interfere
15 significantly with the acquisition of cone beam computerized tomography images.

 The preferred embodiment includes a method for calibration of the radiation therapy delivery system accomplished using a radiation therapy system including the radiation therapy source 409, a collimating structure such as a multi-leaf collimator, and an imager 446. The imager 446 is located opposite the
20 radiation therapy source 409 on a support arm attached to the radiotherapy gantry 406 and in the preferred embodiment is a flat panel imager 404 designed for imaging of the high energy beam 411. The calibration method preferably employs a reference BB 448 embedded in a lucite cube 450 and located at a known location with respect to the isocenter 430 of gantry rotation, as illustrated
25 in FIG. 25. The cube 450 is precisely leveled, and marks on the cube surface project the location of the isocenter within the cube. The reference BB 448 is imaged at angular increments using the radiation therapy source 409 and imager 446 as the gantry 406 rotates through 360°, preferably clockwise and counter-clockwise. In each image, the reference BB 448 is located preferably by an
30 automated centroid calculation, and the edge of each leaf of the multi-leaf collimator and the edge of the collimators are localized by calculation of maximum signal gradient. After subtracting a sinusoid of specified amplitude

from the measured deflections, the residuals represent imperfections in leaf placement. These residuals can then be incorporated into the controller of the multi-leaf collimator and calibrated out. An alternative approach is to modify the planning system to generate "corrected" leaf positions. After calibration, the patient positioning lasers in the treatment room are adjusted to the set of laser alignment marks located on the lucite cube.

The preferred embodiment furthermore includes a calibrator that calibrates the cone beam computerized tomography imaging geometry relative to that of the radiation therapy source 409. Calibration of the cone beam calibration tomography geometry is preferably performed immediately following multi-leaf collimator leaf calibration, without moving the reference BB 448. The same procedure is performed using the x-ray source 402 and the flat panel imager 404; however, in this case, the residuals are used to adjust the back-projection trajectories in the reconstruction process. The image of the localized BB 448 is preferably analyzed using a 3-D centroid algorithm, and the location of isocenter 430 is calculated as a simple offset from the centroid. The isocenter 430 can thus be explicitly identified within the 3-D matrix of cone beam computerized tomography images.

In the preferred embodiment, the cross-calibration of the cone beam computerized tomography imaging system 400 and the radiation therapy delivery system can be tested with a mechanism (phantom) for combined geometry and dosimetry measurement. The phantom preferably includes a water-filled or water-equivalent volume in which a dosimetry insert is rigidly placed at various locations. The dosimetry insert preferably contains either: 1.) a detector matrix of electronic dosimeters, or 2.) a volume of radiosensitive gel dosimeter. In the former case, the dosimeters are embedded in a water-equivalent insert and placed asymmetrically to allow unambiguous identification in a computerized tomography image; furthermore, each dosimeter is sufficiently small as to have legible influence on the dosimetry of other detectors. The electronic signals from the dosimeter matrix are preferably used in either of two ways: 1.) the dosimetry of a complete delivery can be tested by recording the signal from all detectors and comparing to calculations, thereby providing a point dose verification of the

delivery as well as routine pre-treatment quality assurance; and/or 2.) the precision and accuracy of the combined imaging and delivery system can be measured by recording the dose to the detectors as the geometric edge of a leaf can be inferred and compared to the planning system dose calculation. This test is preferably performed for all the leaves in the system by moving the location of the dosimetry insert within the volume. In the case of a radiosensitive gel dosimeter, measurement of 3-D dose distributions delivered by a given treatment scheme can be quantitatively evaluated.

The preferred embodiment furthermore includes delineating the target volume immediately following acquisition of the cone beam computerized tomography image of the patient 441 on the treatment table 443 in the treatment position. Localization of the target volume/lesion 444 and/or surrounding structures can be performed manually, e.g., by contouring of structures in some combination of transaxial, sagittal, coronal, and/or oblique slices. Alternatively, the target volume/lesion 444 and/or surrounding structures can be delineated by an automated localization algorithm, as recognized in the art. In this approach, the target volume/lesion 444 defined in the planning image is overlaid on a given on-line cone beam computerized tomography image, and the images are matched, e.g., by translating and rotating the reference target contour in such a way as to minimize the standard deviation of pixel values compared to the planning image. In the planning image, bony structures are defined, and matching of the planning image with the on-line cone beam computerized tomography image (both with calibrated isocenter positions) on bony structures determines the setup error (rotation and translation) of the bony anatomy. The motion of the soft-tissue target relative to the bony anatomy is quantified by translating and rotating the target volume contours until they cover a homogeneous area (i.e., standard deviation in pixel value differences is minimized).

The treatment plan for the current session can be modified based on the cone beam computerized tomography image data by a number of methods or combinations therein, including recalculation of the RTTP, selection of a modified RTTP from a previously calculated set of plans, and/or

translation, rotation, and/or angulation of the patient. The method chosen should provide a modified plan for the current treatment session in a manner that does not cause uncertainty in the location/orientation of the lesion; therefore, the method should be completed within a short time frame in order to minimize intrafraction organ motion effects, and should not significantly distort patient anatomy. Recalculation of the RTTP based on the cone beam computerized tomography image data should be consistent with such time constraints. Similarly translation, rotation, and/or angulation of the patient should not perturb patient anatomy compared to that measured in the cone beam computerized tomography image, e.g., due to the effects of gravity.

The preferred embodiment entails a streamlined process for rapid lesion localization, selection of an appropriate RTTP, dosimetry review, and transfer of the prescription to the radiation therapy delivery system. The process for on-line cone beam computerized tomography guidance of radiation therapy procedures is illustrated in FIG. 26, which conceptually separates the system into: 1.) the off-line treatment process; 2.) priors for on-line selection and correction; and 3.) the on-line imaging and treatment process.

The off-line treatment process in the preferred embodiment begins with a planning image on which contours of the target volume and surrounding structures are defined, and margins for target deformation, delivery precision, and delineation precision are applied. Inverse planning is performed according to a given protocol for radiation therapy of the given treatment site, e.g., a number of radiation therapy beams directed at the patient from various angles, with target dose uniformity and normal tissue volume constraints to match the prescription. In addition to this reference plan, a plurality of additional plans (the constrained plan set) are generated as a function of various translations and/or rotations of the target volume. Plans are preferably generated at small increments of each possible translation and/or rotation (e.g., rotation of the target volume about the y axis).

In the preferred embodiment for on-line plan selection and correction of lesion localization errors, the target volume/lesion and its relationship to bony structure in the planning image are prepared for use as priors, and the

constrained plan set is transferred to the radiation therapy system to verify deliverability prior to the on-line procedure. In the on-line treatment process, the patient 441 is set up on the treatment table 443 in the treatment position, and cone beam computerized tomography images are acquired as described above. The target volume/lesion 444 and surrounding structures are delineated in the cone beam computerized tomography data, thereby identifying the translations and/or rotations of the target volume/lesion 444 relative to the position and orientation in the planning image. As described above, translations may be corrected by translation of the computer-controlled treatment table 443, and rotations may be corrected by selection of an appropriate plan from the constrained plan set. The translation of the lesion 444 observed in the cone beam computerized tomography image relative to the planning image is corrected by translation of the patient 441 on the treatment table 443 in the y and/or z directions, and/or by rotation about the x axis. The orientation of the lesion 444 (i.e., rotations about the y and/or z axes) are corrected by selecting from the previously calculated constrained plan set a modified RTTP that most closely corresponds to the measured rotation of the lesion 444. Meanwhile, radiographic monitoring of the patient 441 can be used to check against intrafraction motion of the patient 441. Furthermore, a cone beam computerized tomography image acquired immediately prior to, during, or following the treatment procedure can be obtained in order to provide accurate representation of the location of patient anatomy during treatment delivery, which can be stored for off-line review, evaluation, and modification of subsequent treatment sessions. Following transferal of the prescription to the delivery system, the treatment plan is executed according to the patient setup and treatment plan determined from the cone beam computerized tomography image.

The foregoing discussion discloses and describes merely exemplary embodiments of the present invention. One skilled in the art will readily recognize from such discussion, and from the accompanying drawings and claims, that various changes, modifications and variations can be made therein without departing from the spirit and scope of the invention as defined

in the following claims. For example, the cone beam computerized tomography system can be adapted to perform animal testing identification, and non-invasive and non-destructive component structural testing.

WE CLAIM:

1. A radiation therapy system comprising:
a radiation source that moves about a path and directs a beam
of radiation towards an object;
5 a cone-beam computer tomography system comprising:
an x-ray source that emits an x-ray beam in a cone-beam
form towards said object;
an amorphous silicon flat-panel imager receiving x-rays
after they pass through the object, said imager providing an image of
10 said object; and
a computer connected to said radiation source and said cone
beam computerized tomography system, wherein said computer receives said
image of said object and based on said image sends a signal to said radiation
source that controls said path of said radiation source.
- 15 2. The radiation therapy system of claim 1, wherein said x-ray
source comprises a kV x-ray source.
3. The radiation therapy system of claim 1, wherein said kV x-ray
source emits x-rays with energies of approximately 100kV.
4. The radiation therapy system of claim 1, wherein said x-ray
20 source comprises a linear accelerator.
5. The radiation therapy system of claim 1, a stage that moves said
object relative to said x-ray source and said amorphous silicon flat-panel
imager.
- 25 6. The radiation therapy system of claim 5, wherein said stage
rotates about an axis of rotation said object relative to said x-ray source and
said amorphous silicon flat-panel imager.

7. The radiation therapy system of claim 2, a stage that moves said object relative to said x-ray source and said amorphous silicon flat-panel imager.

5 8. The radiation therapy system of claim 7, wherein said stage rotates about an axis of rotation said object relative to said x-ray source and said amorphous silicon flat-panel imager.

9. The radiation therapy system of claim 4, a stage that moves said object relative to said x-ray source and said amorphous silicon flat-panel imager.

10 10. The radiation therapy system of claim 9, wherein said stage rotates about an axis of rotation said object relative to said x-ray source and said amorphous silicon flat-panel imager.

11. The radiation therapy system of claim 1, wherein said x-rays from said x-ray source are emitted along a source plane.

15 12. The radiation therapy system of claim 6, wherein said x-rays from said x-ray source are emitted along a source plane that is perpendicular to said axis of rotation.

20 13. The radiation therapy system of claim 10, further comprising an alignment laser that allows visualization of said axis of rotation and said source plane.

14. The radiation therapy system of claim 1, wherein said amorphous silicon flat-panel imager comprises an array of individual detector elements.

25 15. The radiation therapy system of claim 14, wherein said array is a two-dimensional array.

16. The radiation therapy system of claim 14, wherein each of said individual detector elements comprises a-Si:H photodiode.

17. The radiation therapy system of claim 16, wherein each of said individual detector elements further comprises a transistor coupled to said Si:H photodiode.

5 18. The radiation therapy system of claim 1, wherein said computer receives said image from said amorphous silicon flat-panel imager and generates a computer tomography image of said object based on said received image.

19. The radiation therapy system of claim 1, wherein said image is a two dimensional projection image.

10 20. The radiation therapy system of claim 19, wherein said computer receives said two dimensional projection image from said amorphous silicon flat-panel imager and generates a computer tomography image of said object based on said two dimensional projection image.

15 21. The radiation therapy system of claim 1, further comprising a gantry with a first arm and a second arm, wherein said x-ray source is attached to said first arm and said amorphous silicon flat-panel imager is attached to said second arm.

22. The radiation therapy system of claim 21, wherein said gantry rotates about an axis of rotation.

20 23. The radiation therapy system of claim 22, wherein said gantry rotates about a second axis of rotation.

24. The radiation therapy system of claim 21, wherein said gantry is attached to a wall of a room.

25 25. The radiation therapy system of claim 23, wherein said gantry is attached to a wall of a room.

26. The radiation therapy system of claim 22, wherein said gantry is attached to a mobile platform that can translationally move on a floor of a room.

5 27. The radiation therapy system of claim 23, wherein said gantry is attached to a mobile platform that can translationally move on a floor of a room.

10 28. The radiation therapy system of claim 1, wherein said radiation source operates at a power level higher than that of said x-ray source, wherein said radiation is of an intensity and energy that is effective for radiation treatment of an area of said object.

29. The radiation therapy system of claim 21, wherein said radiation source operates at a power level higher than that of said x-ray source, wherein said radiation is of an intensity and energy that is effective for radiation treatment of an area of said object.

15 30. The radiation therapy system of claim 1, wherein said x-ray source is coincident with said radiation source.

31. The radiation therapy system of claim 1, wherein said x-ray source is displaced relative to said radiation source.

20 32. The radiation therapy system of claim 1, wherein operation of said cone beam computerized tomography system with an external trigger that controls a biological process of a patient in which said object is located.

33. The radiation therapy system of claim 32, wherein said external trigger comprises an active breathing control mechanism.

25 34. The radiation therapy system of claim 32, wherein said external trigger comprises a cardiac gating mechanism.

35. The radiation therapy system of claim 1, further comprising an imaging device positioned opposite said radiation source and generating an

image of said object based on radiation from said radiation source that passes through said object.

36. An imaging system comprising:
an x-ray source that emits x-rays towards an object;
5 an imager that receives x-rays from said object based on said emitted x-rays and forms an image of said object;
an imager support system that attaches said imager to a support structure, wherein said imager support system comprises:
a first arm having one end attached to said imager and
10 another end attached to said support structure; and
a second arm having one end attached to said imager and another end attached to said support structure.

37. The imaging system of claim 36, wherein said imager support system comprises a third arm having one end attached to said imager and
15 another end attached to said support structure.

38. The imaging system of claim 37, wherein said third arm lies in a plane that bisects a line segment that joins said one ends of said first and second arms.

39. The imaging system of claim 38, wherein said imager is
20 symmetrically positioned with respect to said plane.

40. The imaging system of claim 38, wherein said imager is asymmetrically positioned with respect to said plane.

41. The imaging system of claim 38, further comprising a motorized system that moves said imager from a position where said imager is
25 symmetrically positioned with respect to said plane to a position where said imager is asymmetrically positioned with respect to said plane.

42. The imaging system of claim 37, wherein each of said another ends of said first, second and third arms are attached to a pivot which is attached to said support structure.

5 43. The imaging system of claim 36, wherein said support structure comprises a rotating drum of a gantry.

44. The imaging system of claim 43, wherein said x-ray source is attached to said rotating drum.

45. The imaging system of claim 44, wherein said x-ray source translates in a direction that is parallel to an axis of rotation of said drum.

10 46. The imaging system of claim 43, further comprising a radiation source attached to said rotating drum.

47. The imaging system of claim 36, wherein said x-rays emitted from said x-ray source are emitted in a cone beam form.

15 48. The imaging system of claim 36, wherein said imager comprises an amorphous silicon flat-panel imager.

49. An imaging system comprising:
an x-ray source that emits x-rays towards an object;
an imager that receives x-rays from said object based on said emitted x-rays and forms an image of said object;
20 an imager support system that attaches said imager to a support structure, wherein said imager support system comprises:
a pivoting arm that has one end pivotably attached to said support structure and another end pivotably attached to said imager.

25 50. The imaging system of claim 49, wherein said support structure comprises a rotating drum of a gantry.

51. The imaging system of claim 50, wherein said x-ray source is attached to said rotating drum.

52. The imaging system of claim 51, wherein said x-ray source translates in a direction that is parallel to an axis of rotation of said drum.

5 53. The imaging system of claim 50, further comprising a radiation source attached to said rotating drum.

54. The imaging system of claim 49, wherein said x-rays emitted from said x-ray source are emitted in a cone beam form.

10 55. The imaging system of claim 49, wherein said imager comprises an amorphous silicon flat-panel imager.

56. An imaging system comprising:
an x-ray source that emits x-rays towards an object;
an imager that receives x-rays from said object based on said emitted x-rays and forms an image of said object;
15 an imager support system that attaches said imager to a support structure, wherein said imager support system comprises:
a C-arm attached to a support structure, wherein said imager is attached to one end of said C-arm.

20 57. The imaging system of claim 56, wherein said C-arm moves along an arc.

58. The imaging system of claim 56, wherein said support structure comprises a rotating drum of a gantry.

59. The imaging system of claim 58, wherein said x-ray source is attached to another end of said C-arm.

25 60. The imaging system of claim 58, further comprising a radiation source attached to said rotating drum.

61. The imaging system of claim 56, wherein said x-rays emitted from said x-ray source are emitted in a cone beam form.

62. The imaging system of claim 56, wherein said imager comprises an amorphous silicon flat-panel imager.

5 63. A method of treating an object with radiation, comprising:
move a radiation source about a path;
direct a beam of radiation from said radiation source towards an
object;
emitting an x-ray beam in a cone beam form towards an object;
10 detecting x-rays that pass through said object due to said
emitting an x-ray beam with an amorphous silicon flat-panel imager;
generating an image of said object from said detected x-rays;
and
controlling said path of said radiation source based on said
15 image.

64. The method of claim 63, wherein x-rays within said x-ray beam have an energy of approximately 100kV.

20 65. The method of claim 63, comprising rotating about an axis of rotation said object relative to said x-ray source and said amorphous silicon flat-panel imager.

66. The method of claim 63, wherein said amorphous silicon flat-panel imager comprises an array of individual detector elements.

67. The method of claim 66, wherein said array is a two-dimensional array.

25 68. The method of claim 66, wherein each of said individual detector elements comprises a-Si:H photodiode.

69. The method of claim 66, wherein said generating comprises forming a computer tomography image of said object based on said detected x-rays.

5 70. The method of claim 65, further comprising rotating about a second axis of rotation said object relative to said x-ray source and said amorphous silicon flat-panel imager.

10 71. The method of claim 63, further comprising emitting a second set of x-rays, separate from said x-rays emitted from said x-ray source, that have an intensity and energy that is effective for radiation treatment of an area of said body.

72. The method of claim 71, wherein said second set of x-rays has an intensity and energy greater than said x-rays emitted from said x-ray source.

15 73. The method of claim 69, further comprising correcting for offset and gain prior to said generating.

74. The method of claim 63, wherein said object comprises an animal.

75. The method of claim 63, wherein said image delineates soft tissue within said animal.

20 76. The method of claim 75, wherein said soft tissue is selected from the group consisting of fat, a muscle, a kidney, a stomach, a bowel and a liver.

25 77. The method of claim 65, wherein said image is formed after one rotation of said body relative to said x-ray source and said amorphous silicon flat-panel imager.

78. The method of claim 63, wherein said x-ray beam is generated by an x-ray source that moves independently of said amorphous silicon flat-

panel imager, said x-ray source moves on a sinusoidal or sawtooth path constrained to a surface of a cylinder while said amorphous silicon panel imager moves in a circular path on a surface of a cylinder.

5 79. The method of claim 78, further comprising adjusting a collimator in real time to adjust a shape of said x-ray beam so it is confined to an active area of said amorphous silicon flat panel imager.

10 80. The method of claim 63, wherein said x-ray beam is generated by an x-ray source that moves dependently of said amorphous silicon flat-panel imager, said x-ray source and said amorphous silicon flat-panel imager each moves on a sinusoidal trajectory on a spherical surface.

 81. A method of adding an auxiliary imaging system to an existing radiation therapy system, said method comprising:
 providing an existing radiation therapy system that comprises a radiation source that is supported on a support structure; and
15 attaching an imager that does not directly face said radiation source to said support structure.

 82. The method of claim 81, wherein said attaching comprises:
attaching said imager to an imager support system;
 forming an opening in said support structure;
20 inserting a male member through an opening formed in said imager support system and said opening formed in said support structure; and
 attaching said inserted male member to said support structure and said imager support system.

25 83. The method of claim 82, wherein said attaching said inserted male member comprises tightening a nut onto said male member.

 84. The method of claim 81, wherein said support structure comprises a rotating drum.

85. The method of claim 82, wherein said support structure comprises a rotating drum.

86. The method of claim 81, further comprising attaching an x-ray source to said support structure.

5 87. The method of claim 84, further comprising attaching an x-ray source to said rotating drum.

88. The method of claim 85, further comprising attaching an x-ray source to said rotating drum.

10 89. A method of delineating a target volume located within a body and shown in a computerized tomography image, comprising:
forming a computerized tomography image of a target volume in a body; and
manually localizing said target volume in a slice of said computerized tomography image.

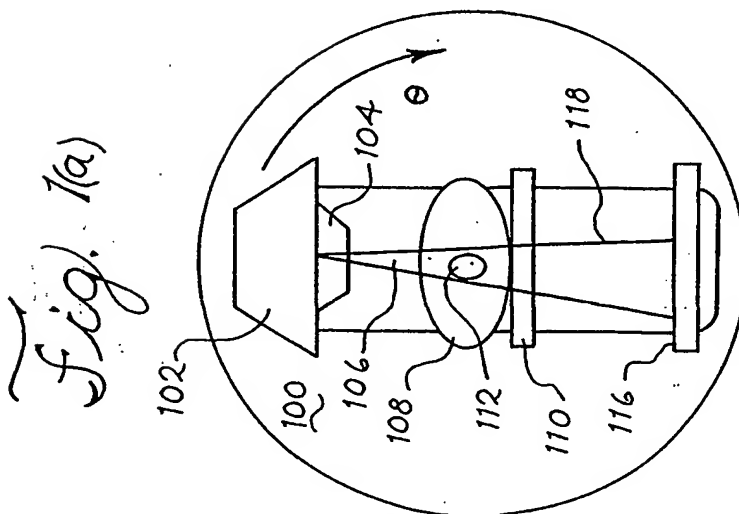
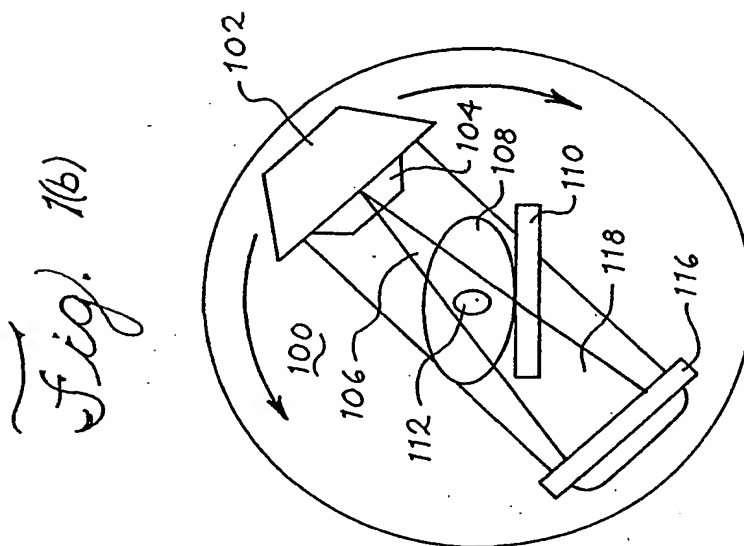
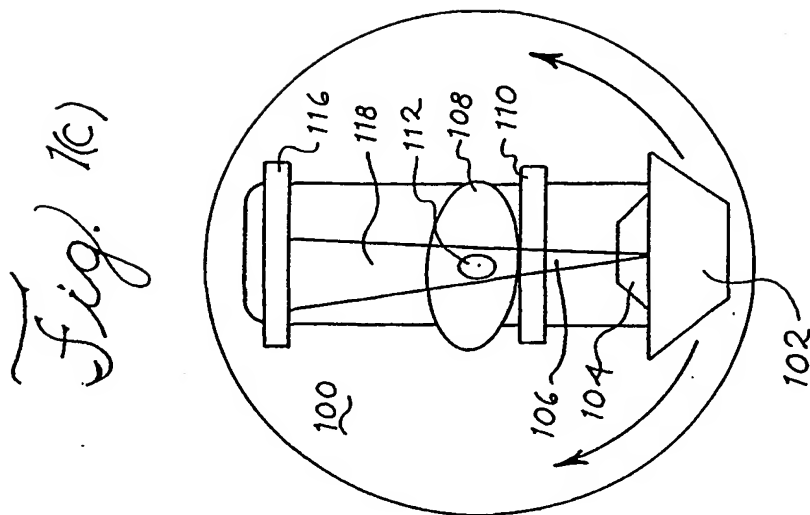
15 90. The method of claim 89, wherein said manually localizing comprises contouring of structures in said slice.

91. The method of claim 89, wherein said manually localizing comprises contouring of structures of said slice in combination with one or more different slices of said computerized tomography image.

20 92. A method of delineating a target volume located within a body and shown in a computerized tomography image, comprising:
forming a computerized tomography image of a target volume in a body;
detecting said target volume
25 identifying translations and/or rotations of said target volume relative to a position and orientation in a planning image;
adjusting the location and/or orientation of said target volume;

selecting a radiation therapy treatment plan from a previously formed set of radiation therapy treatment plans as a function of said detected location and/or orientation of said target volume.

- 5 93. A method of delineating a target volume located within a body and shown in a computerized tomography image, comprising:
- forming a computerized tomography image of a target volume in a body;
- identifying translations and/or rotations of said target volume relative to a position and orientation in a planning image;
- 10 adjusting the location and/or orientation of said target volume;
- calculating a radiation therapy treatment plan based on said adjusted location and/or orientation of said target volume.



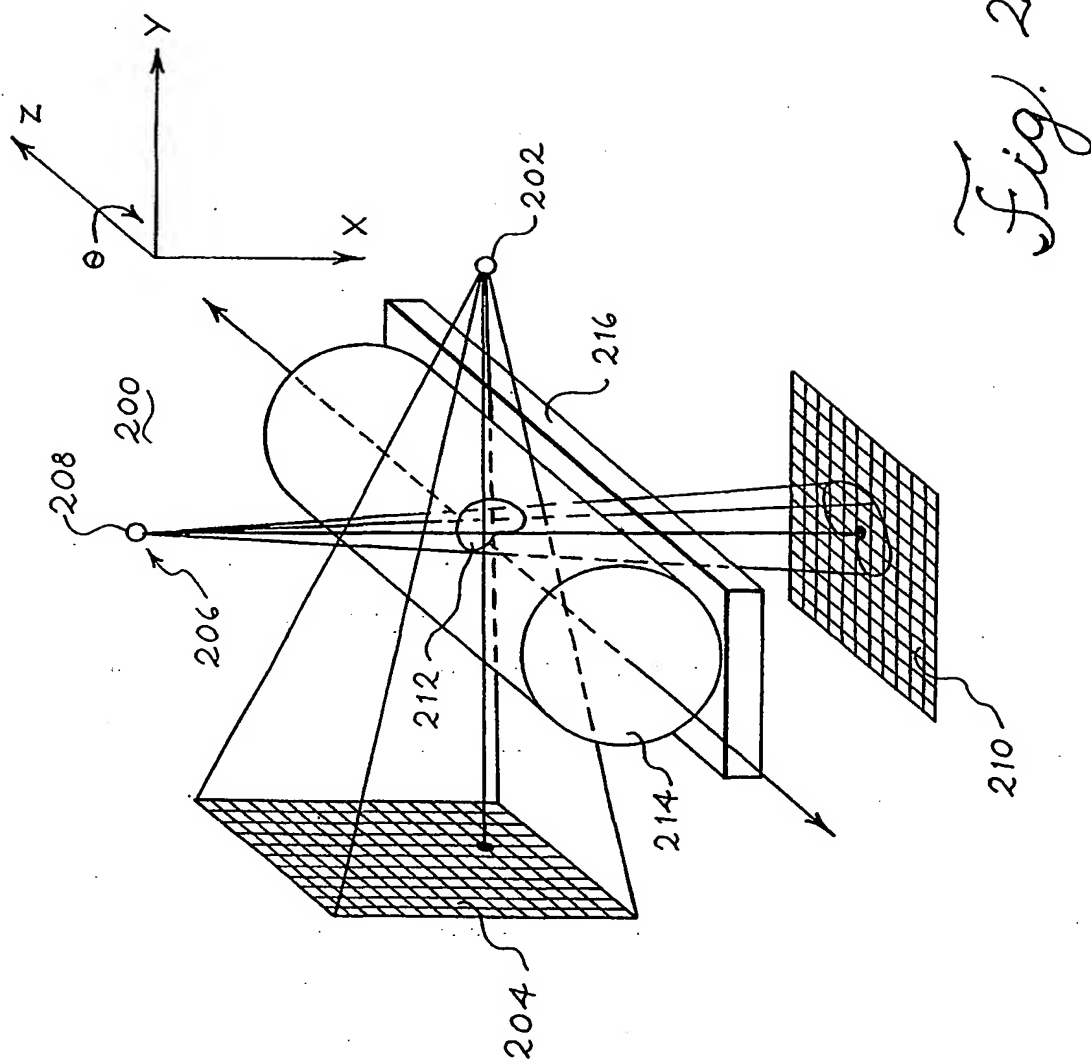
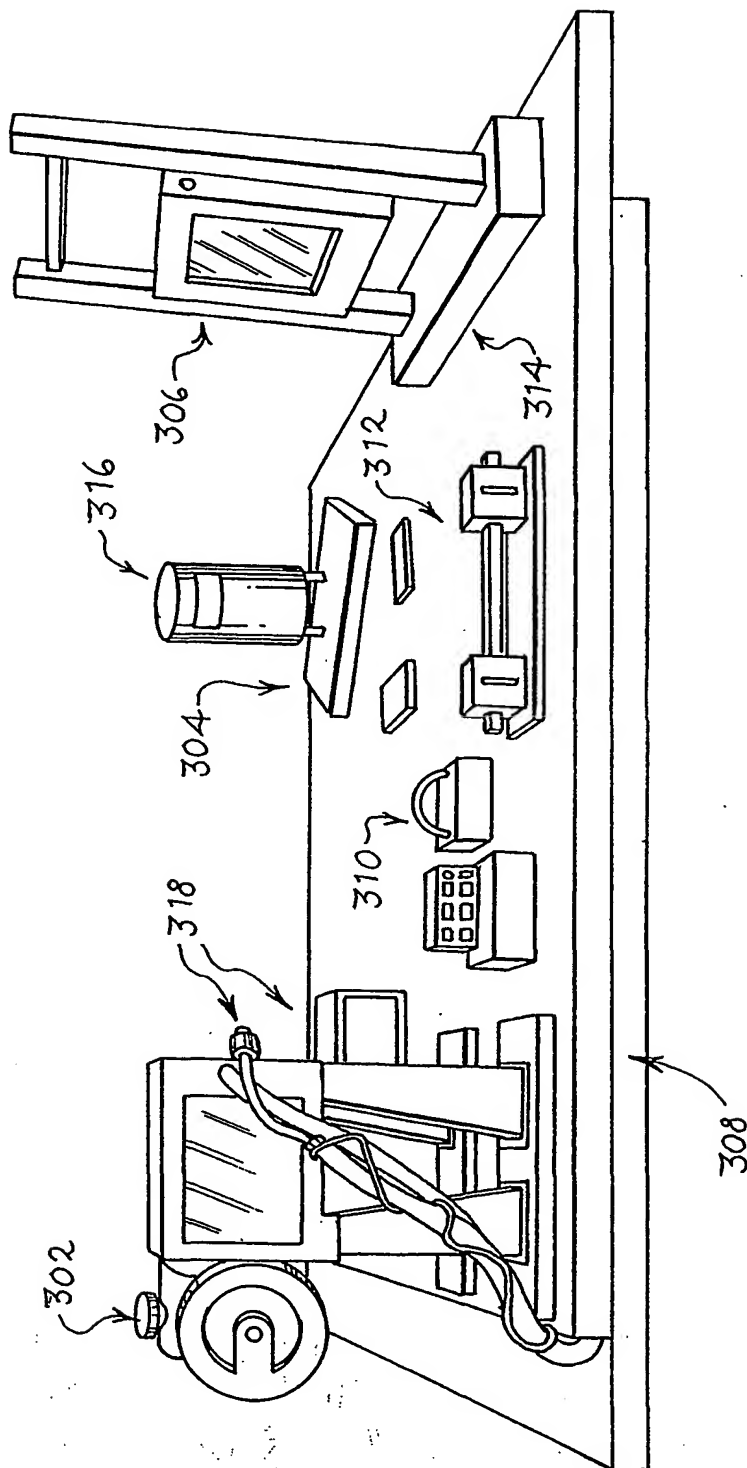
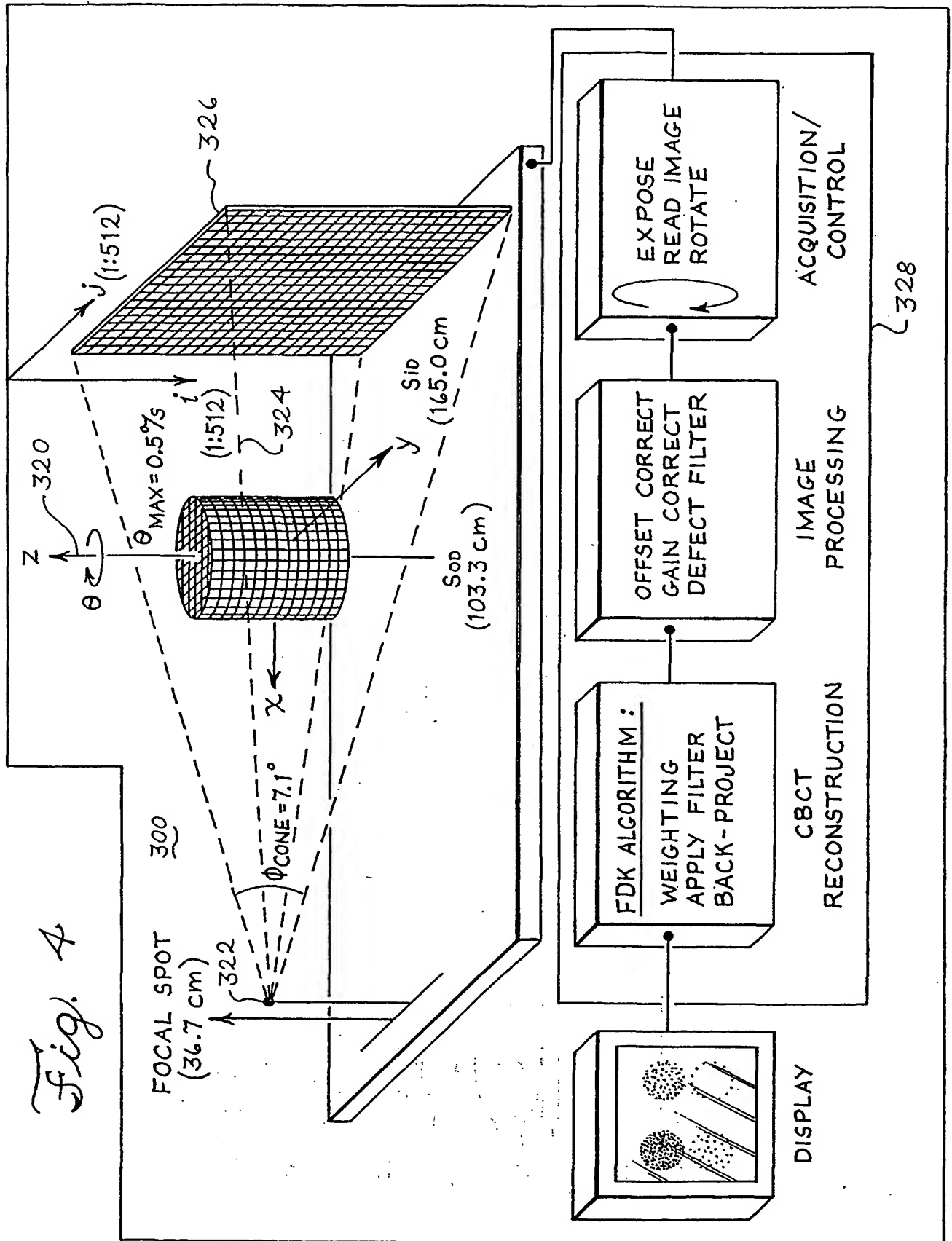
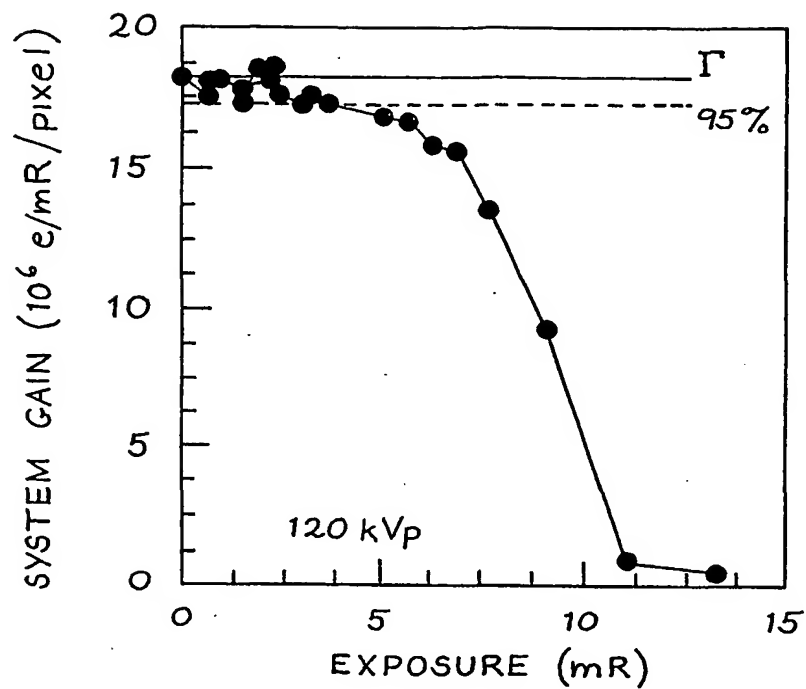
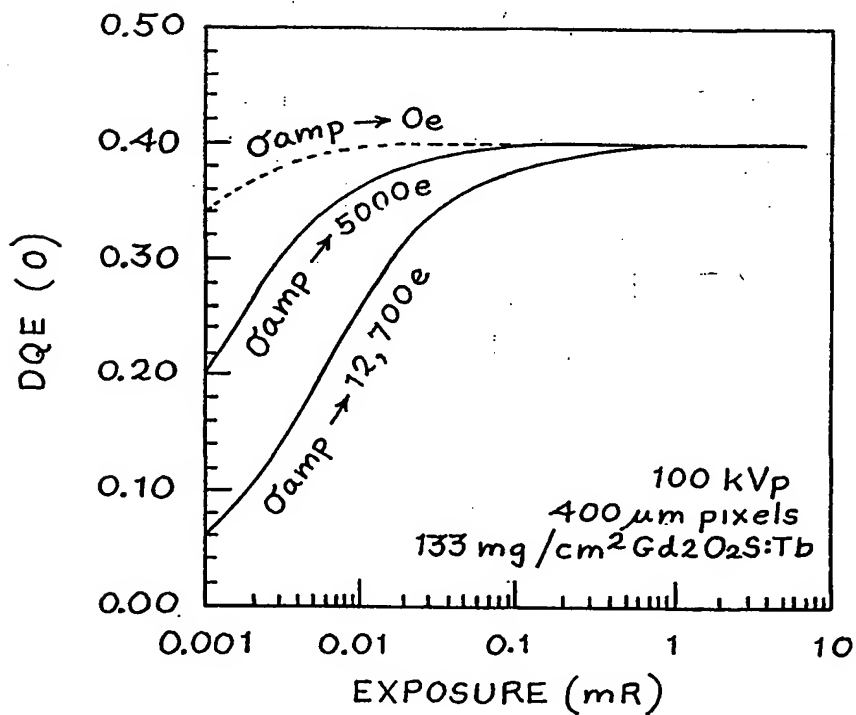
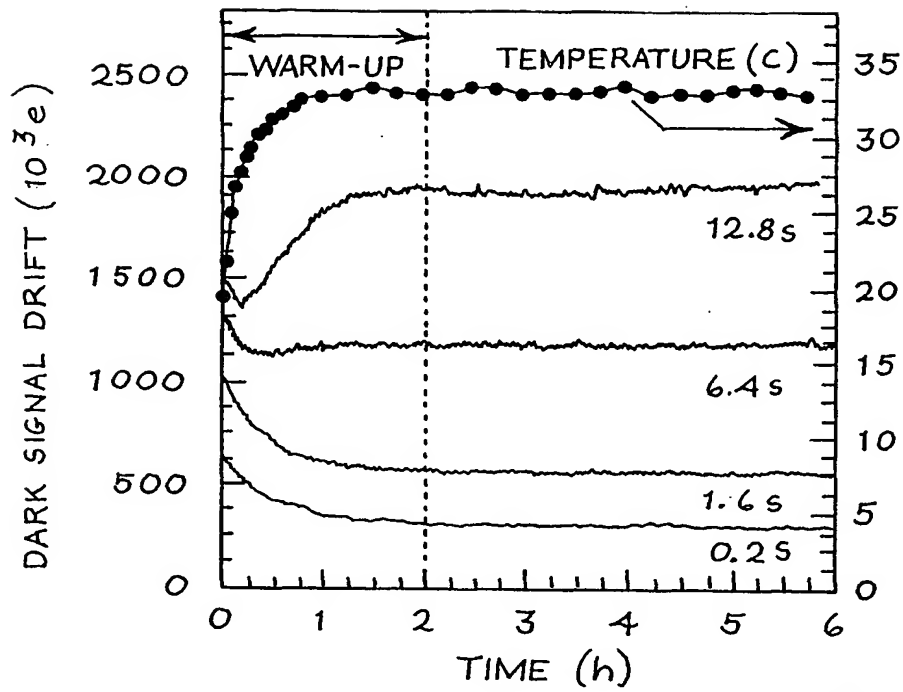
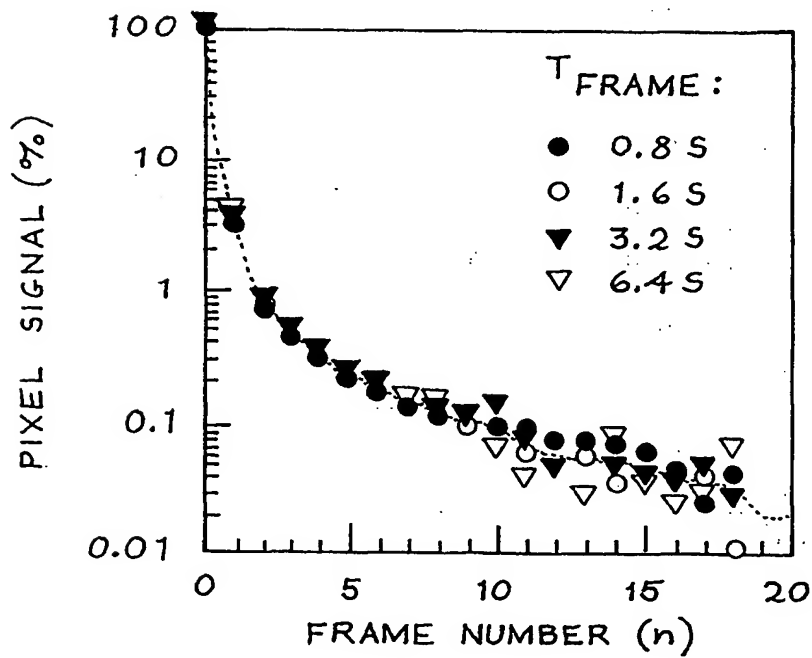


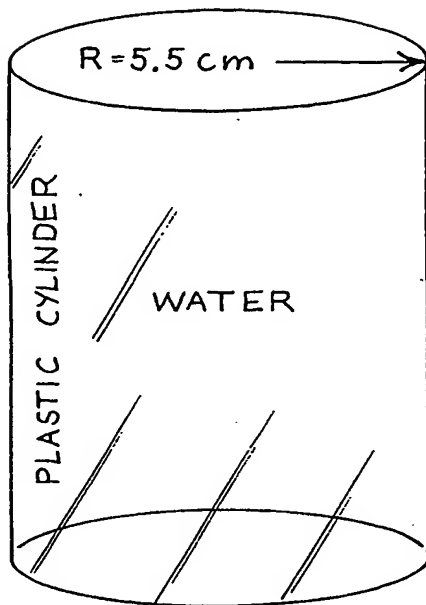
Fig. 3





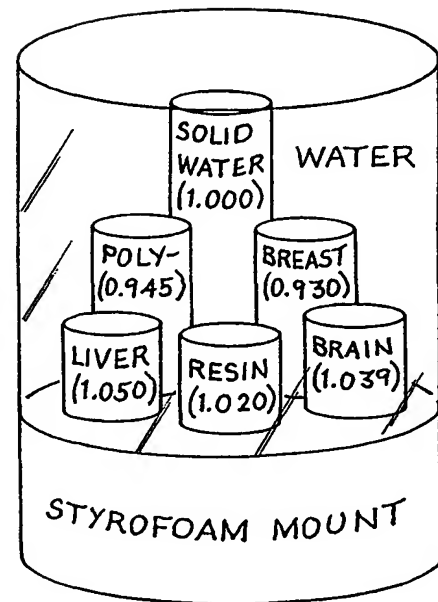
*Fig. 5(a)**Fig. 5(b)*

*Fig. 5(c)**Fig. 5(d)*



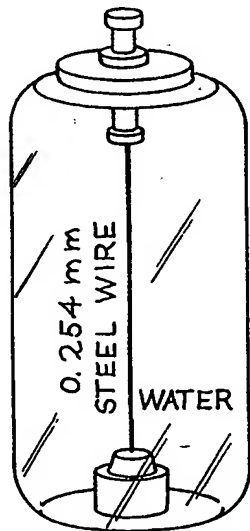
WATER CYLINDER
(UNIFORMITY, NOISE, NPS)

Fig. 6(a)



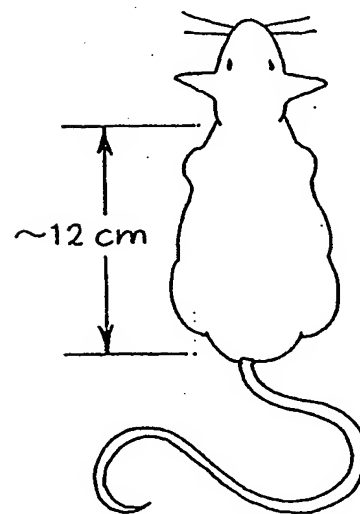
ELECTRON DENSITY PHANTOM
(CONTRAST, CT LINEARITY)

Fig. 6(b)



WIRE PHANTOM
(SPATIAL RESOLUTION, MTF)

Fig. 6(c)



EUTHENIZED WISTAR RAT
(SOFT TISSUE VISIBILITY, BONY DETAIL)

Fig. 6(d)

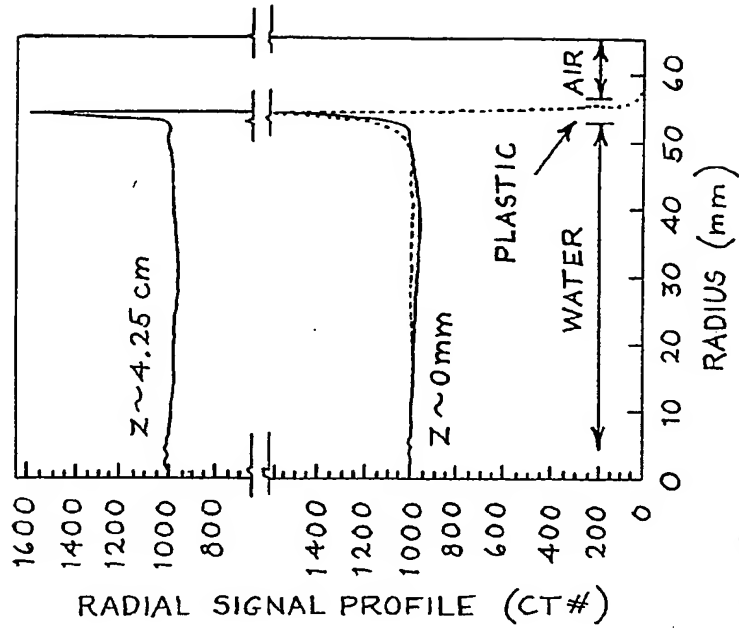
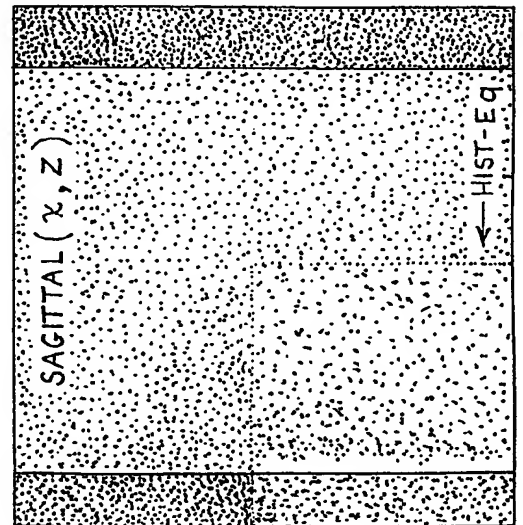
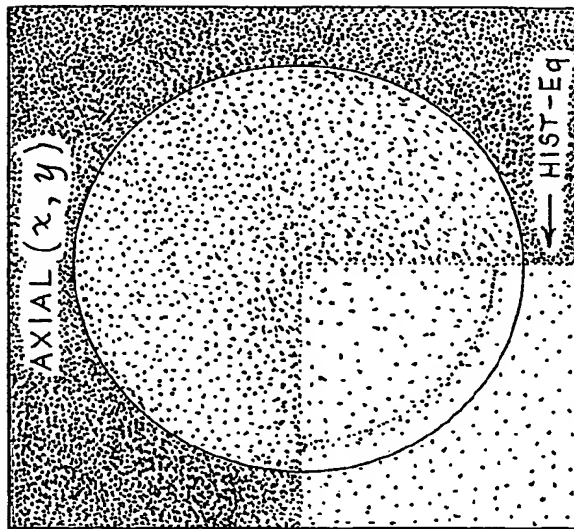
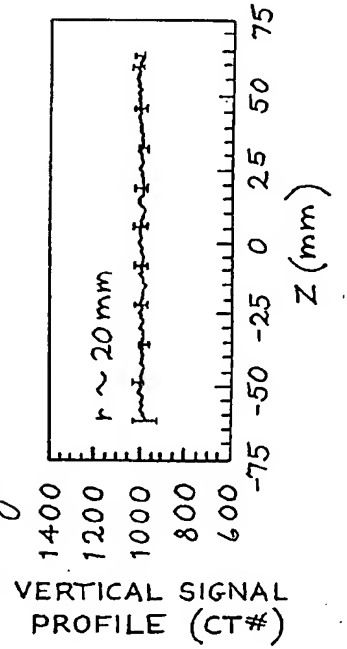


Fig. 7(d)



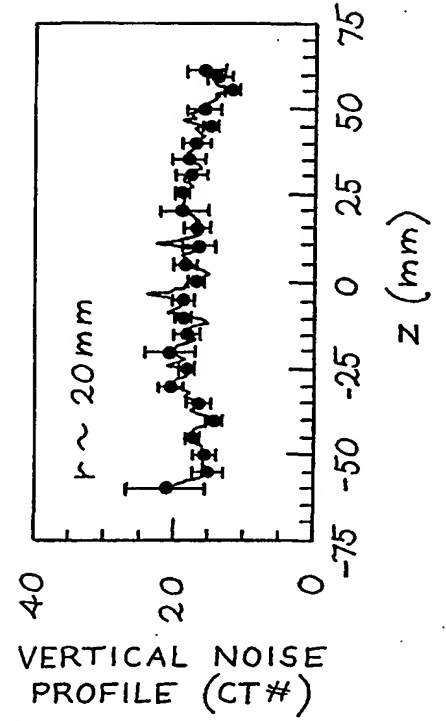
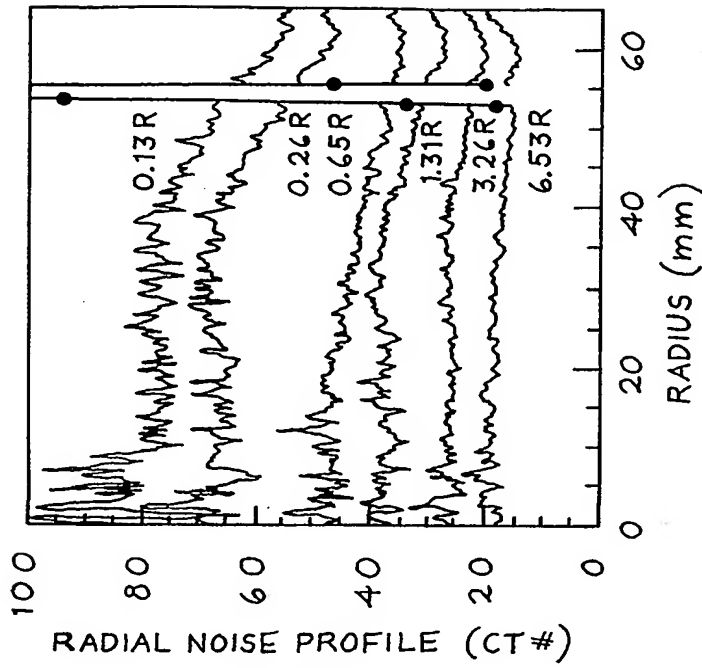
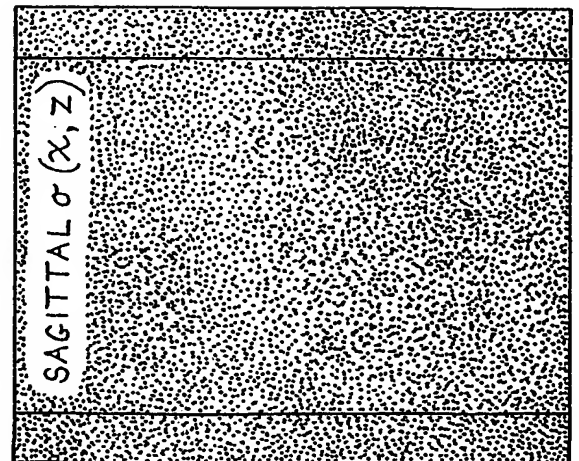
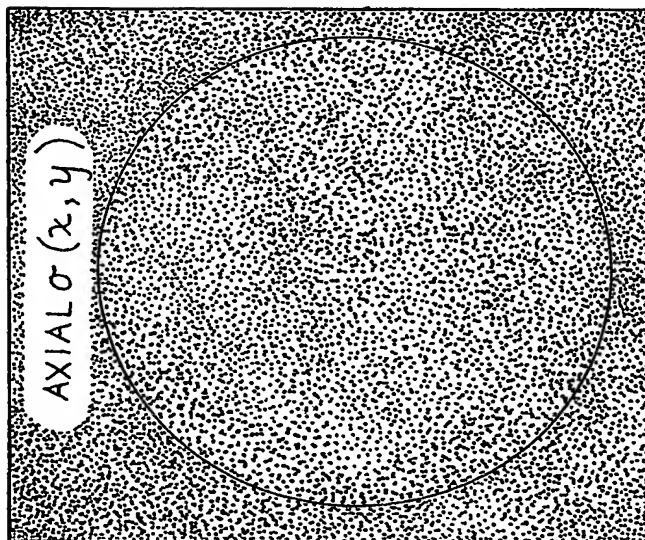
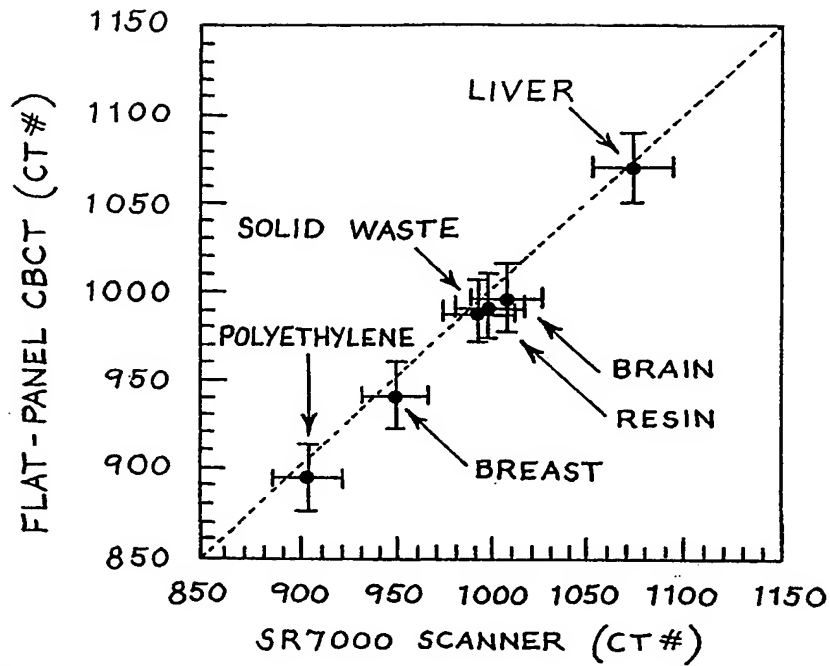
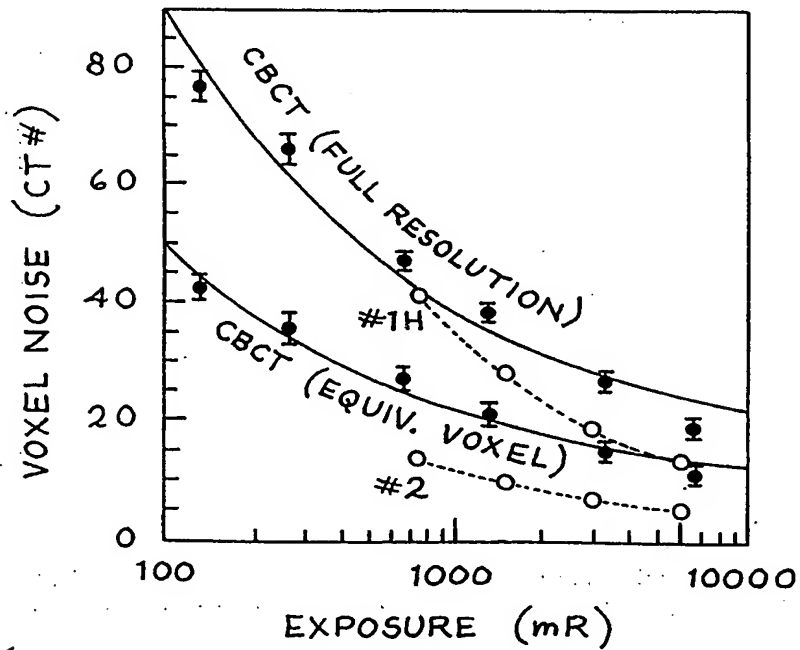


Fig. 8(c)

Fig 8(a)

Fig. 8(d)

Fig. 8(b)

*Fig. 9(a)**Fig. 9(b)*

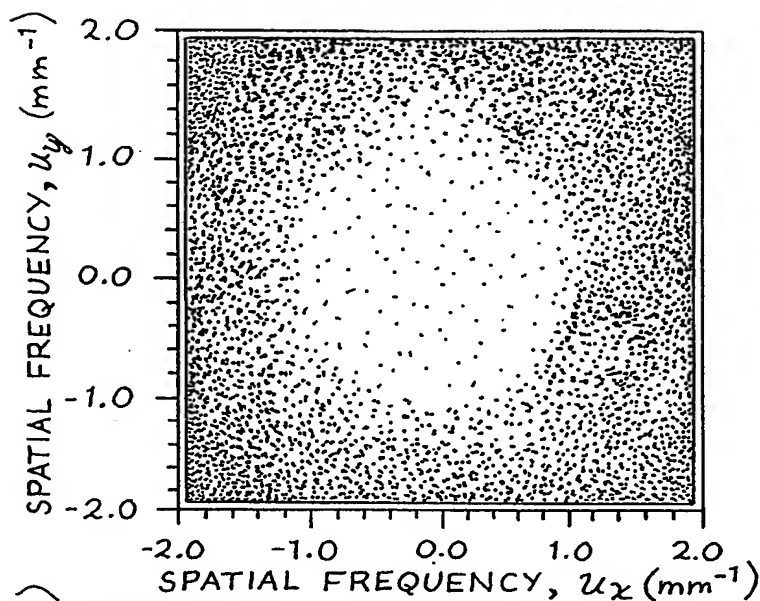
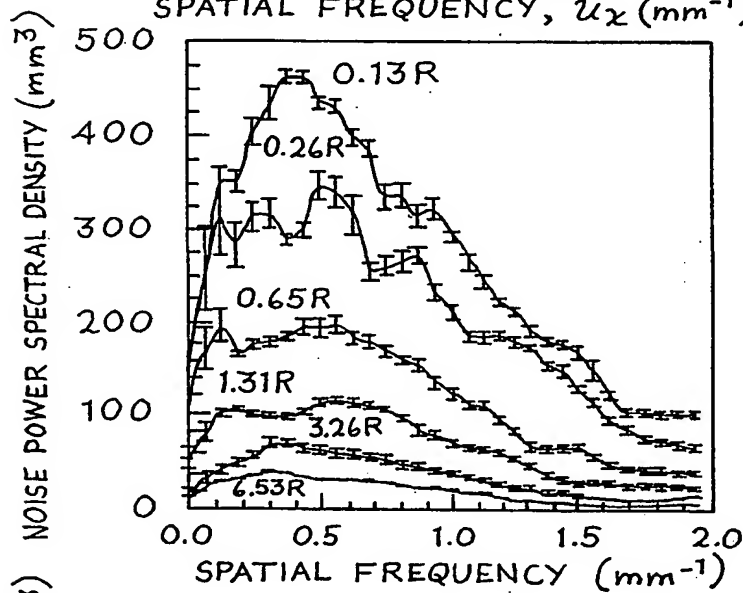
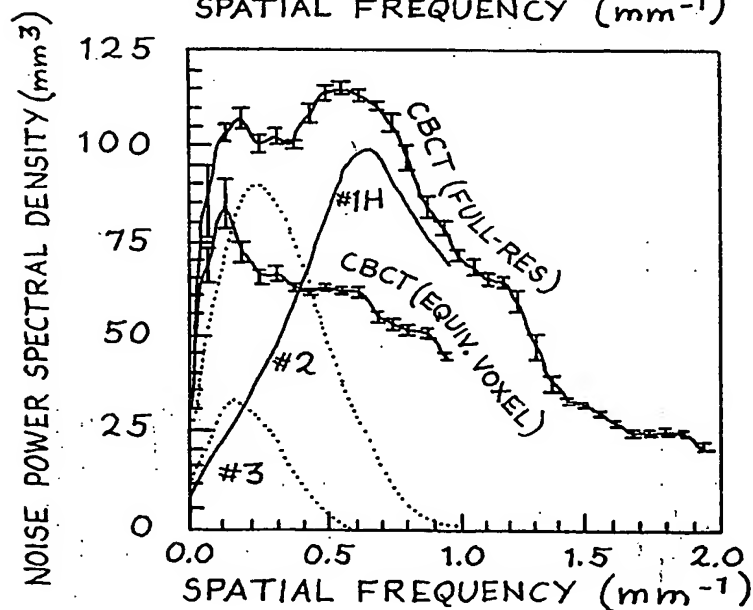
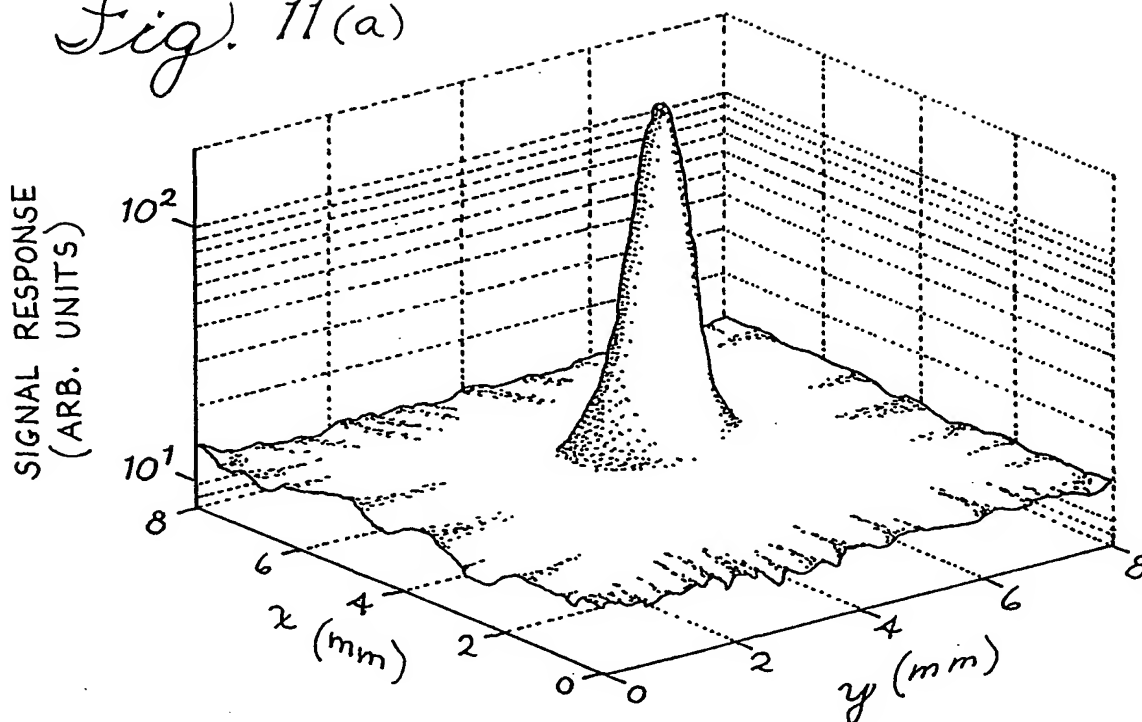
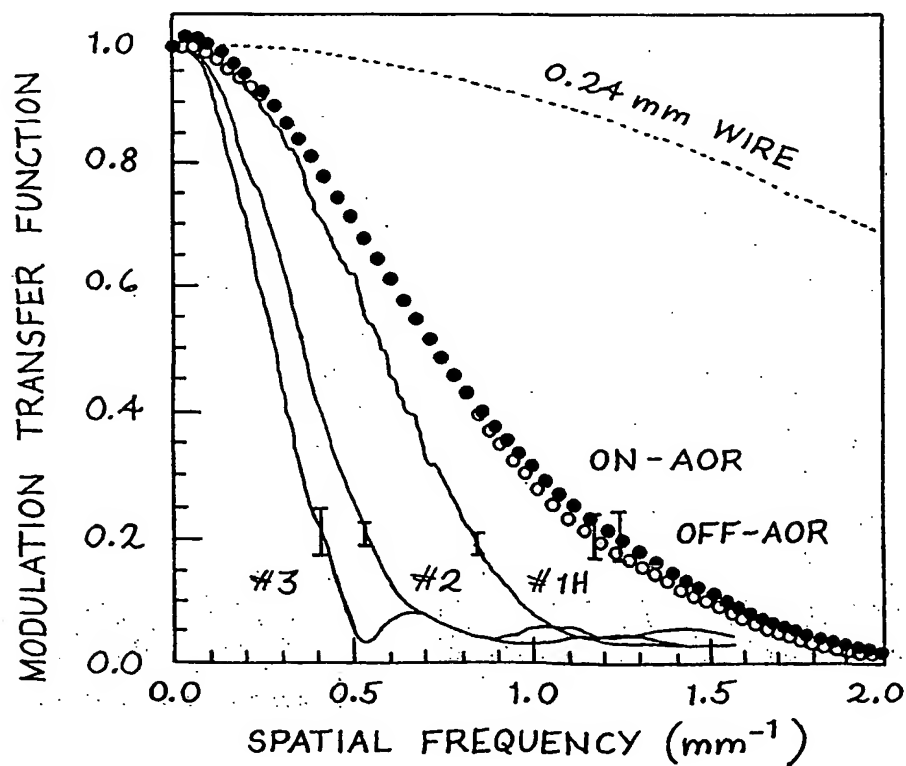
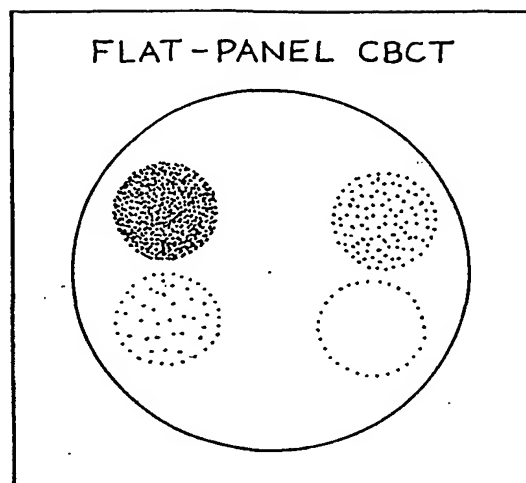
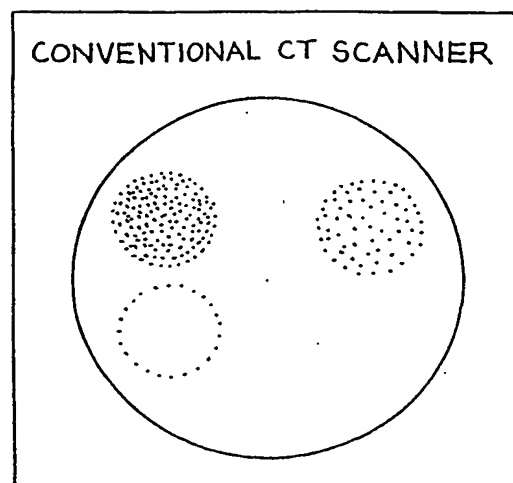
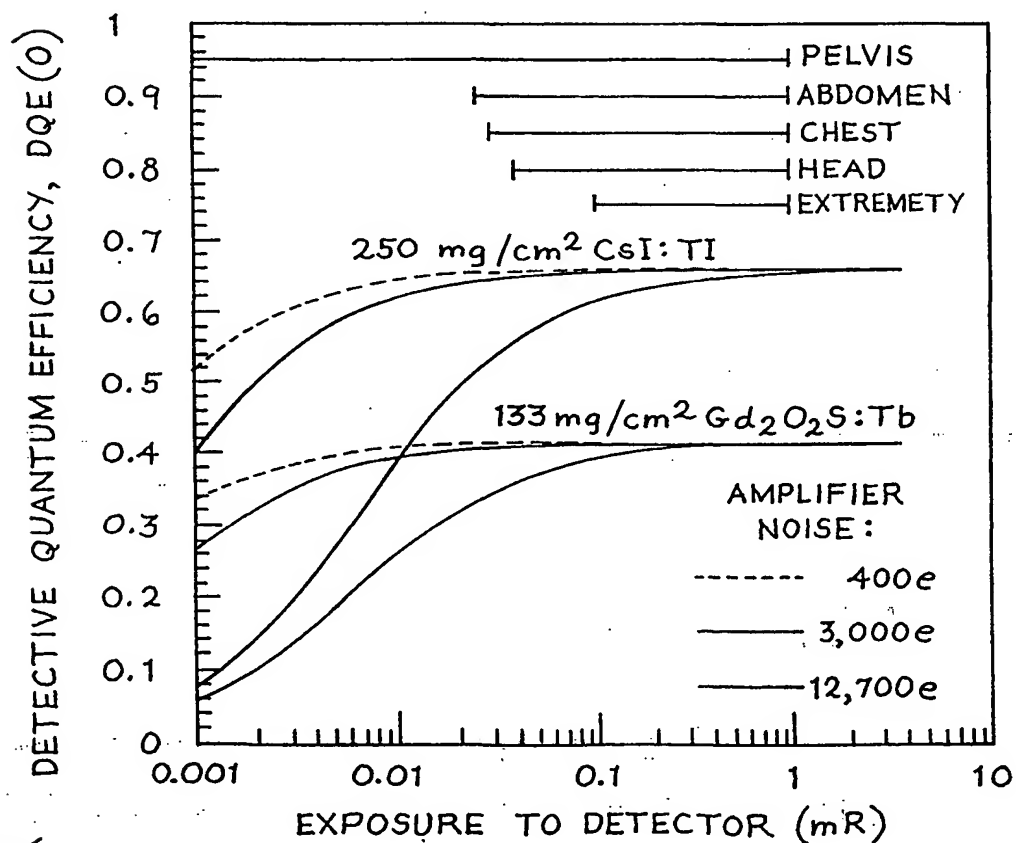
*Fig. 10(a)**Fig. 10(b)**Fig. 10(c)*

Fig. 11(a)*Fig. 11(b)*

*Fig. 12 (a)**Fig. 12 (b)**Fig. 16*

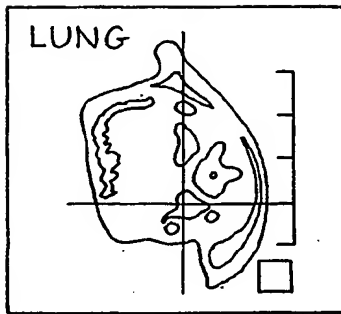


Fig. 13(a)

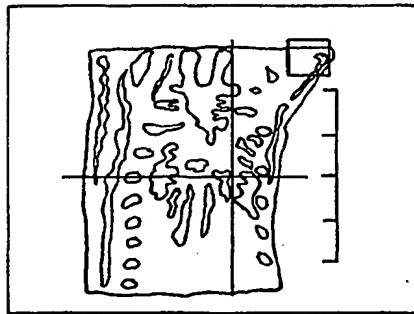


Fig. 13(b)

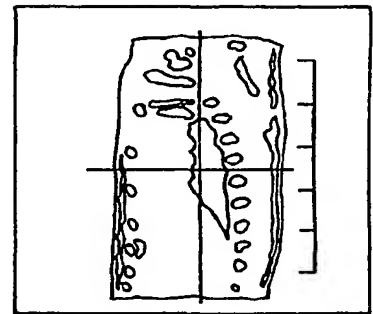


Fig. 13(c)

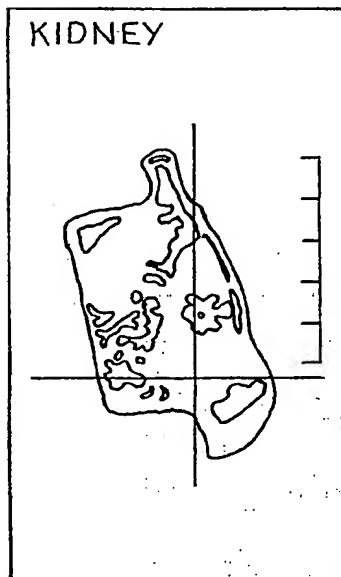


Fig. 13(d)

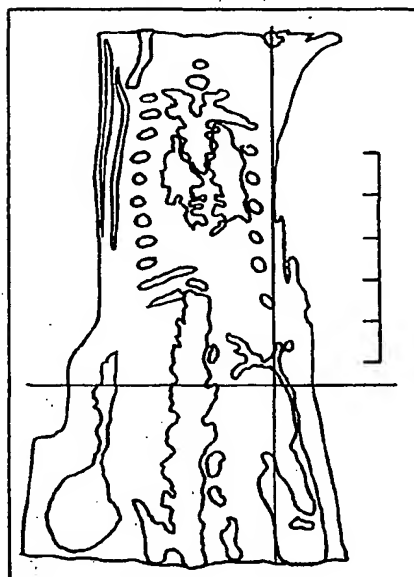


Fig. 13(e)

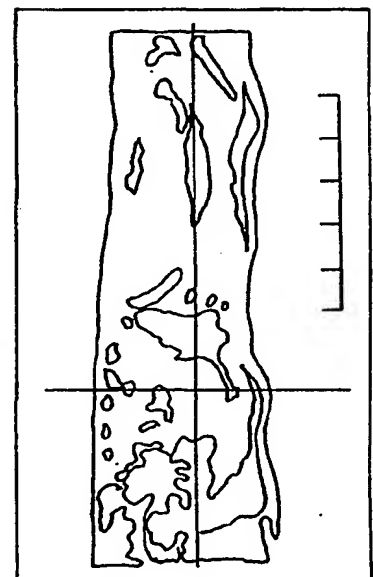


Fig. 13(f)

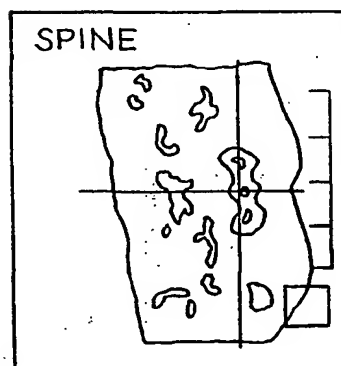


Fig. 13(g)

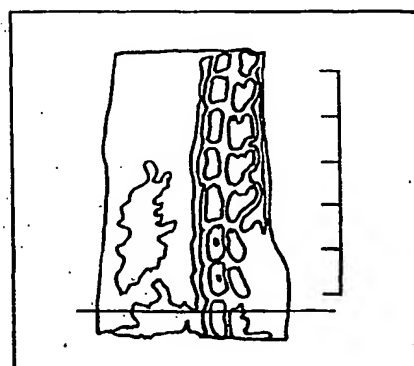


Fig. 13(h)

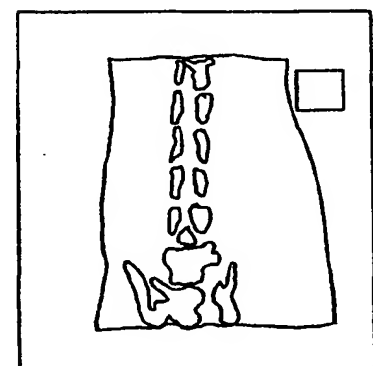


Fig. 13(i)

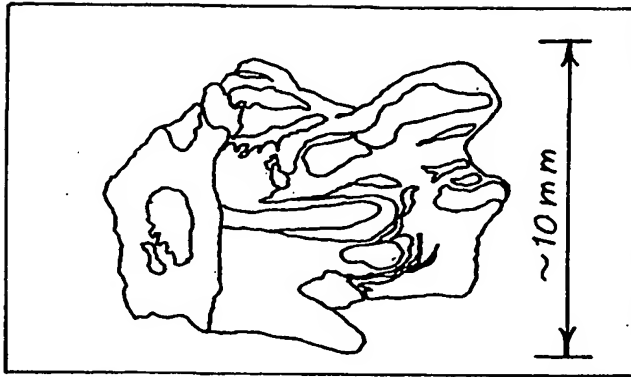


Fig. 14(d)

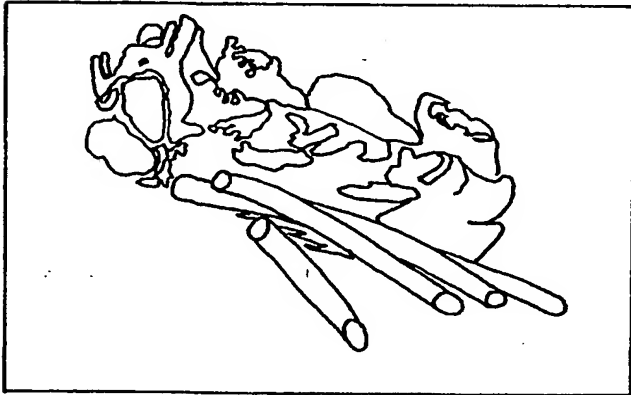


Fig. 14(c)

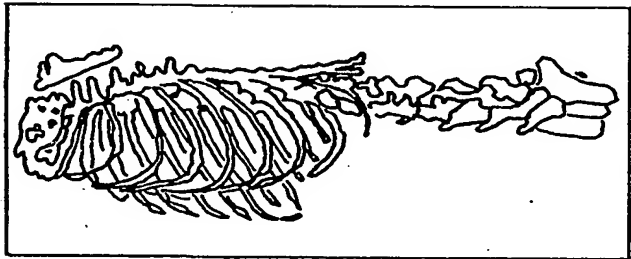


Fig. 14(b)

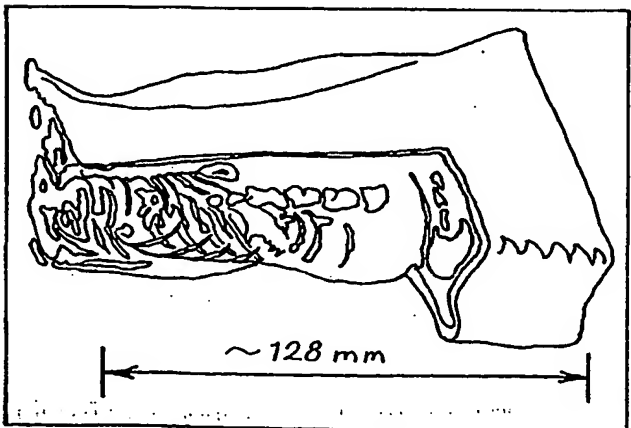


Fig. 14(a)

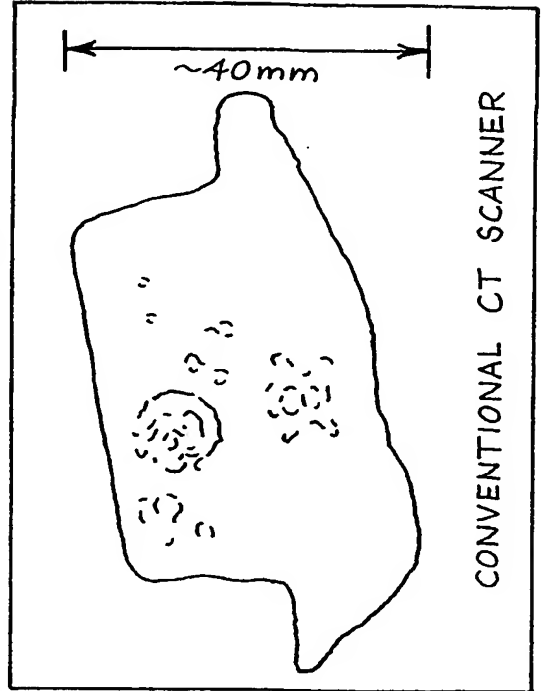


Fig. 15(a)

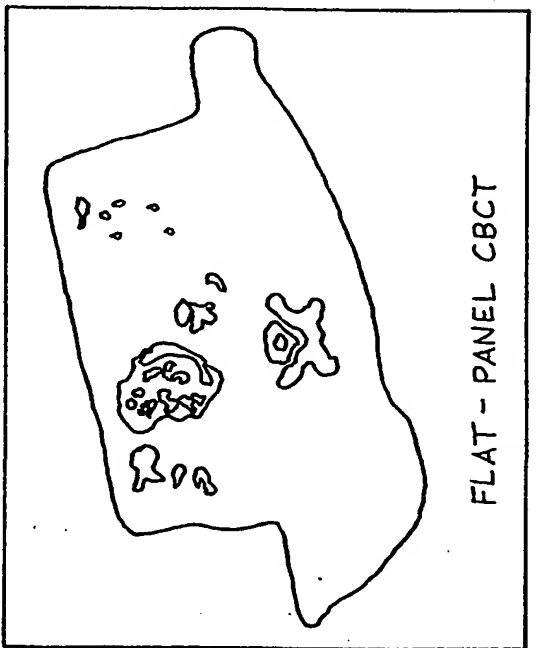


Fig. 15(b)

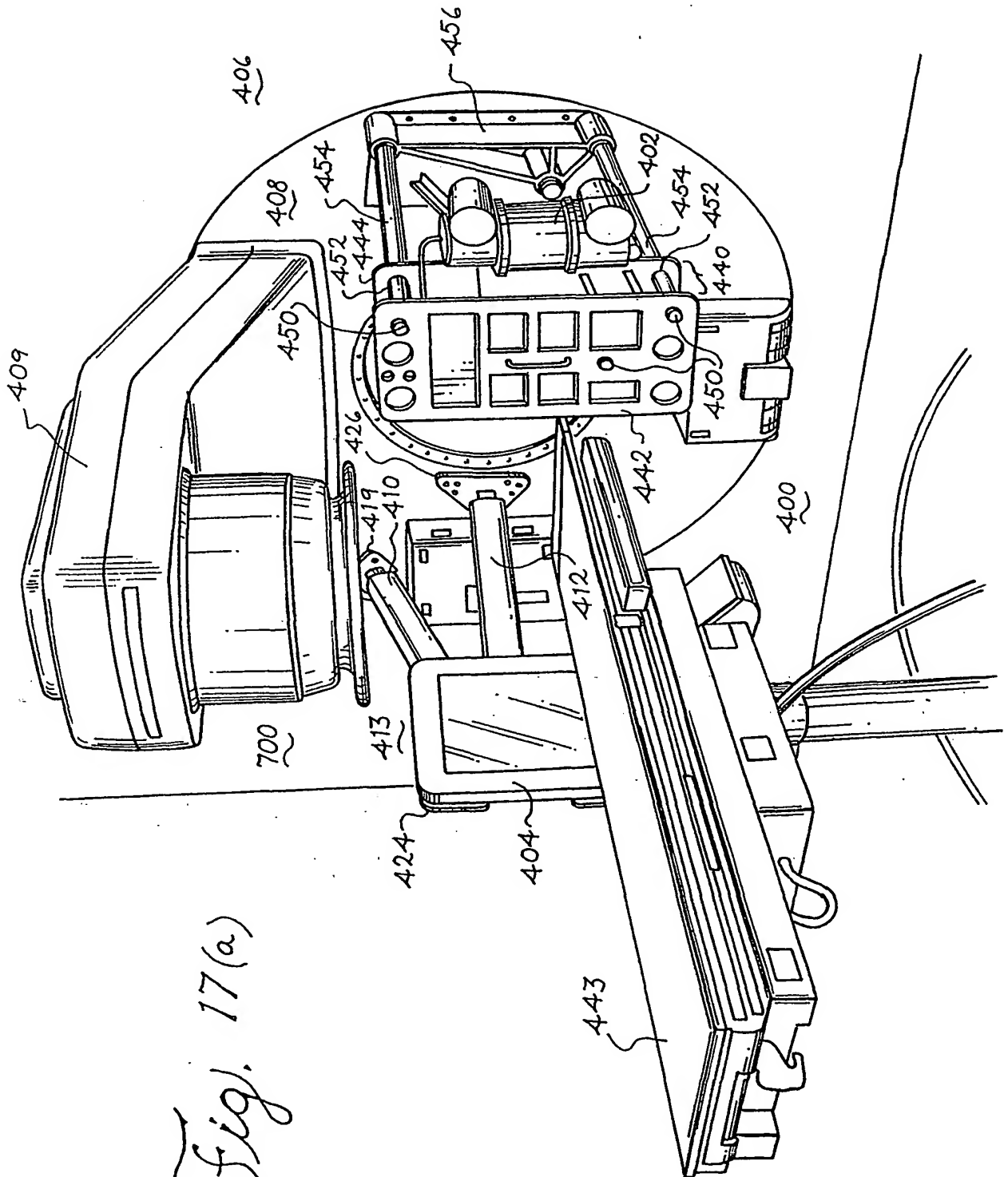
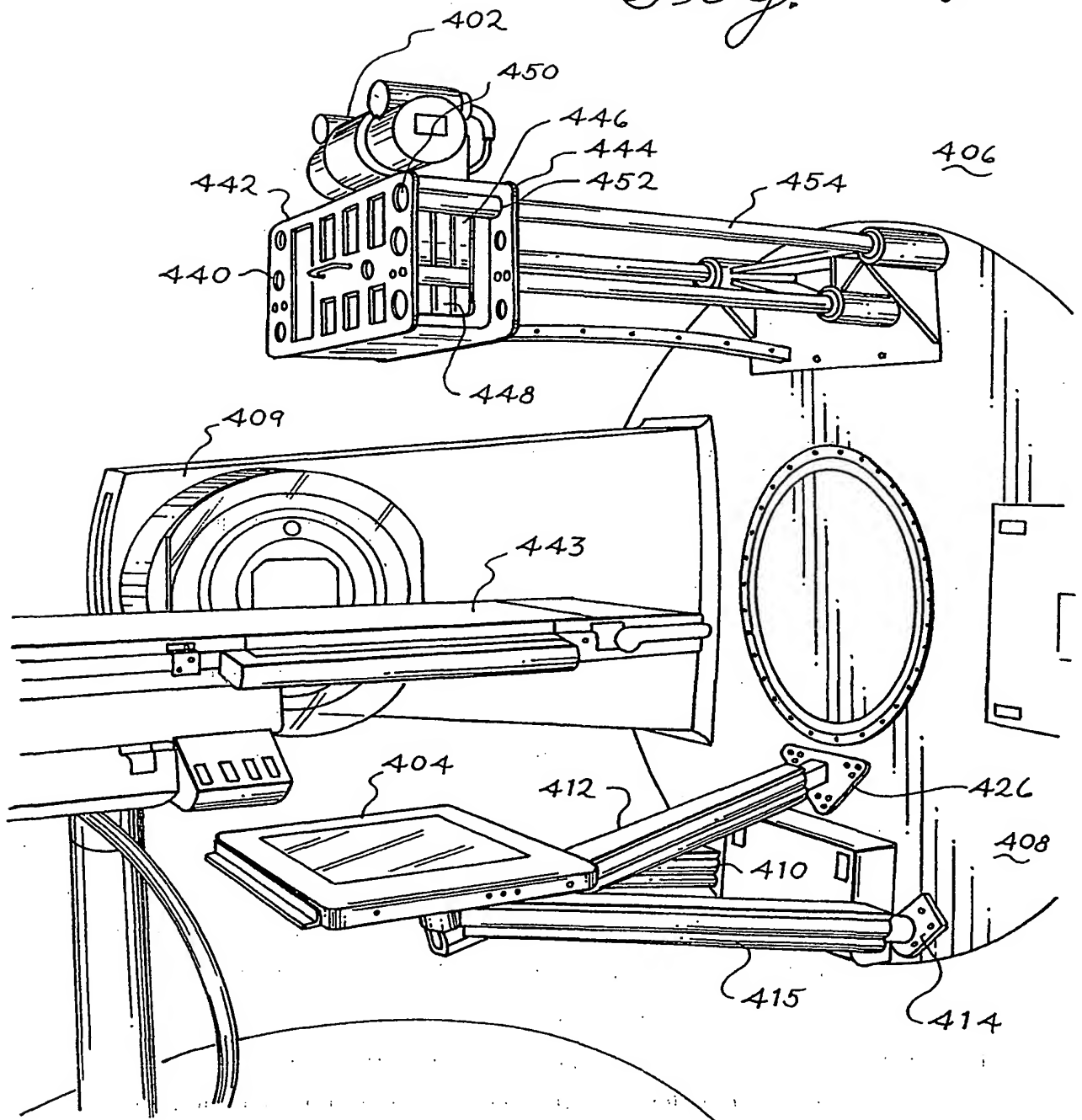


Fig. 17(a)

Fig. 17(b)

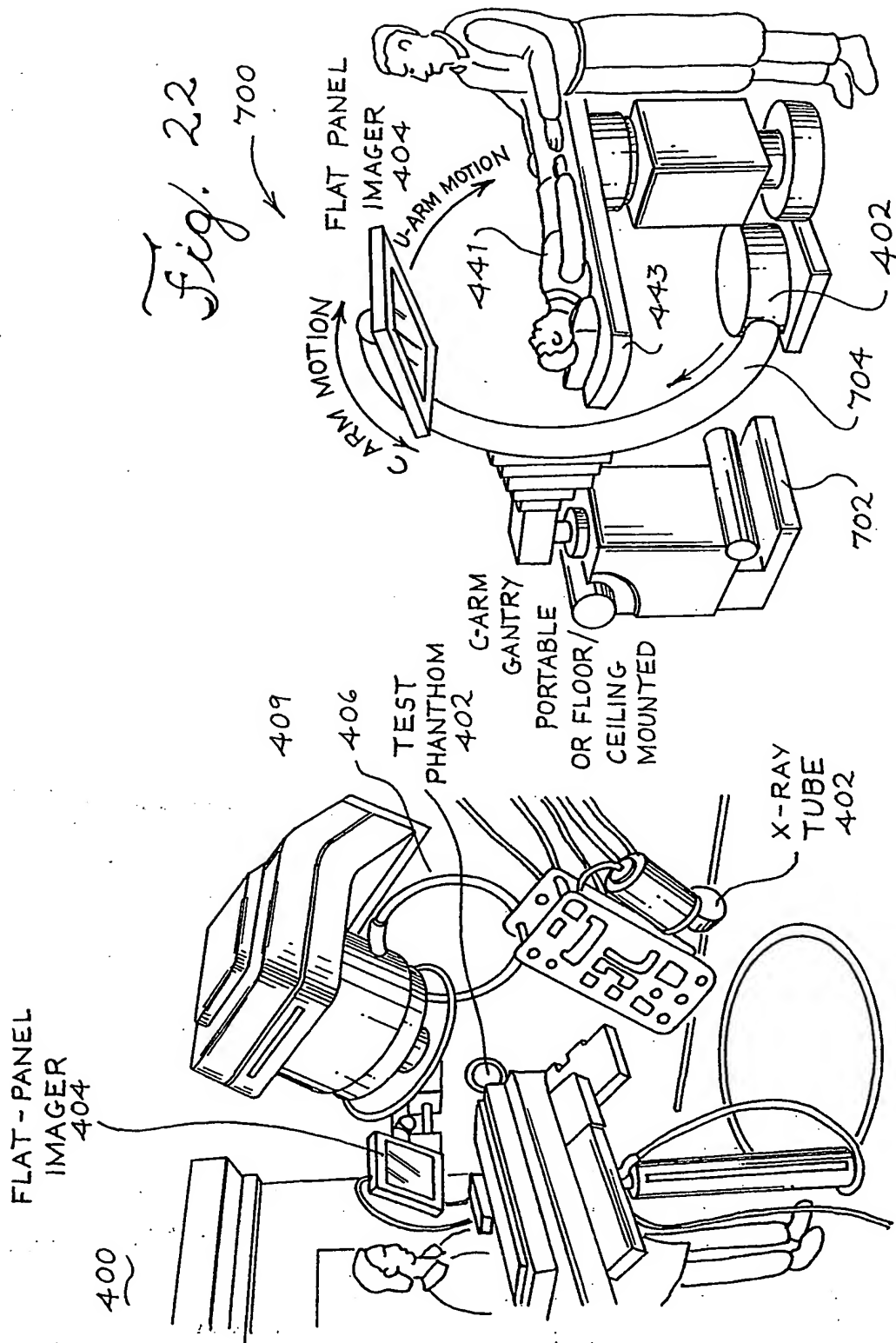


Fig. 17(c)

Fig. 17(d)

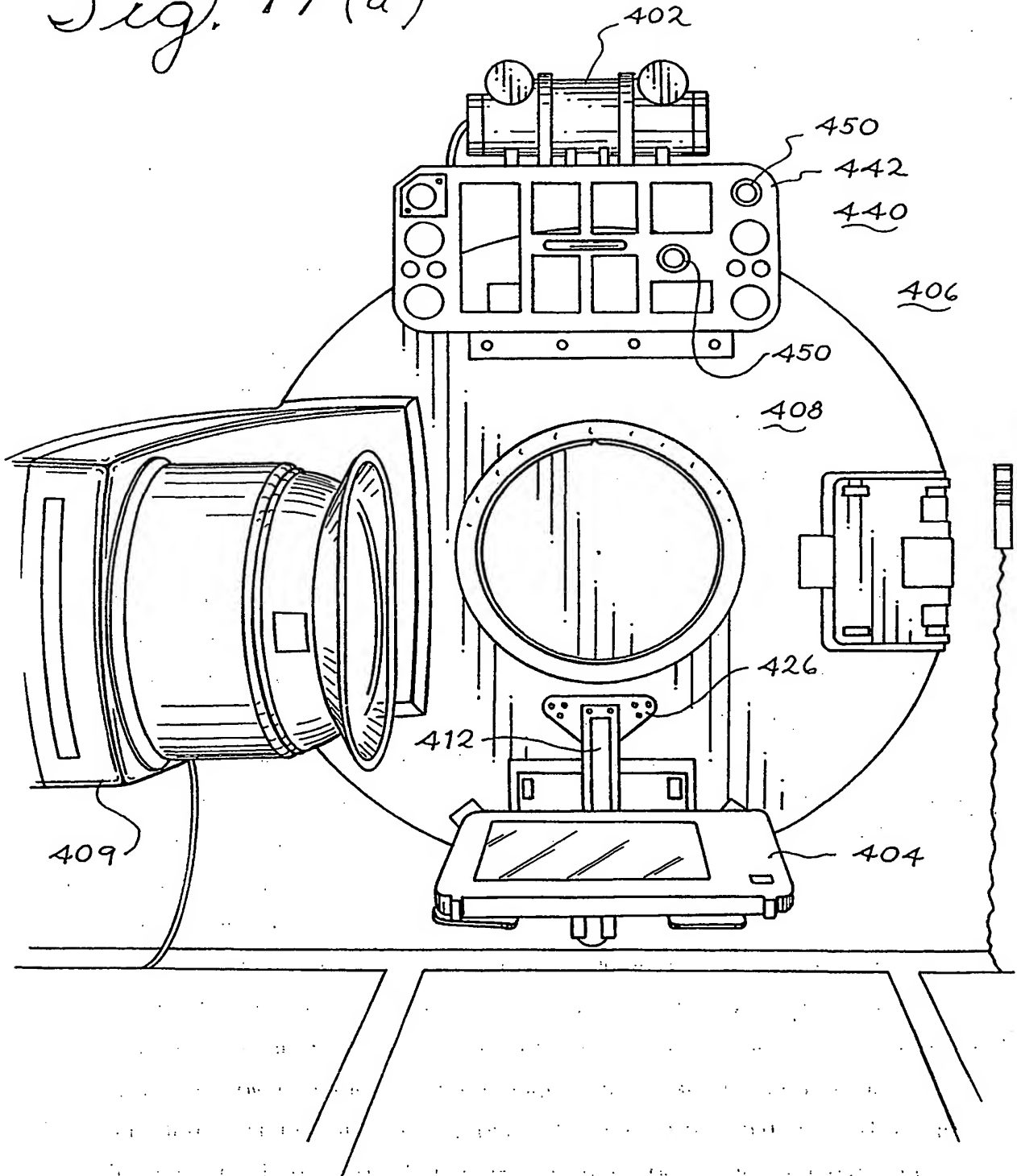


Fig. 17 (e)

Fig. 19(a)

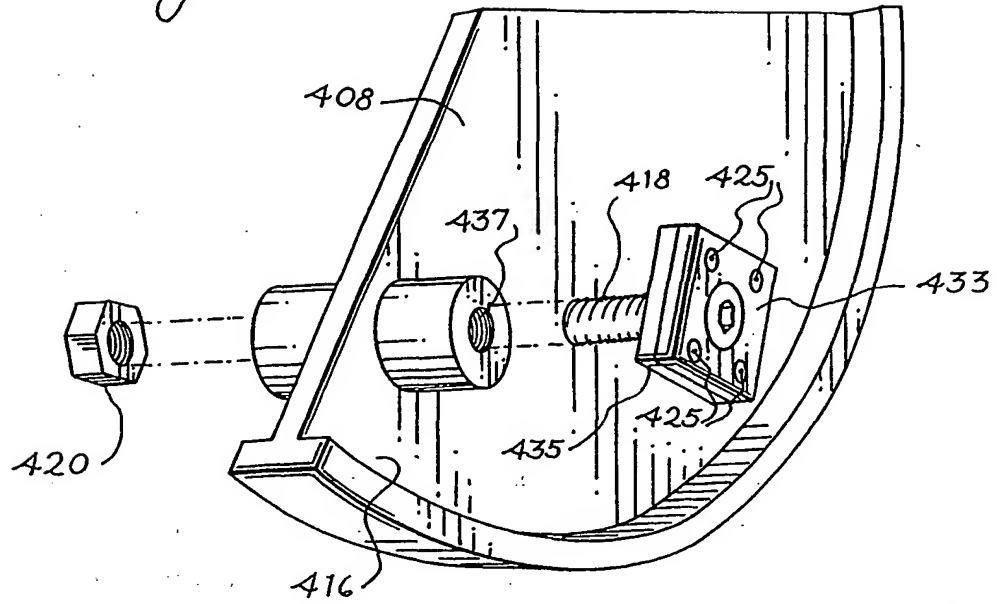
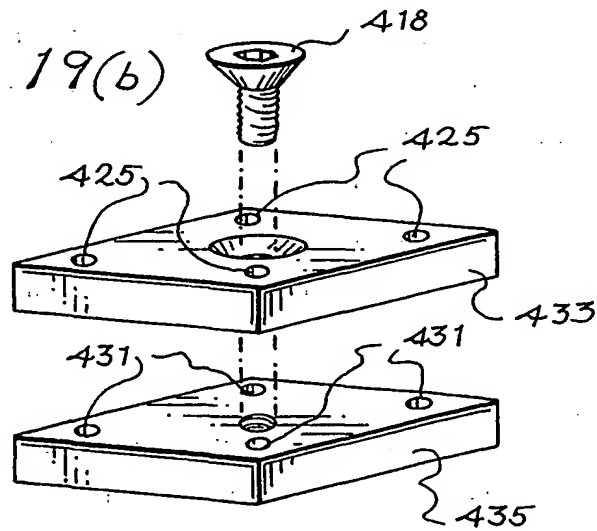


Fig. 19(b)



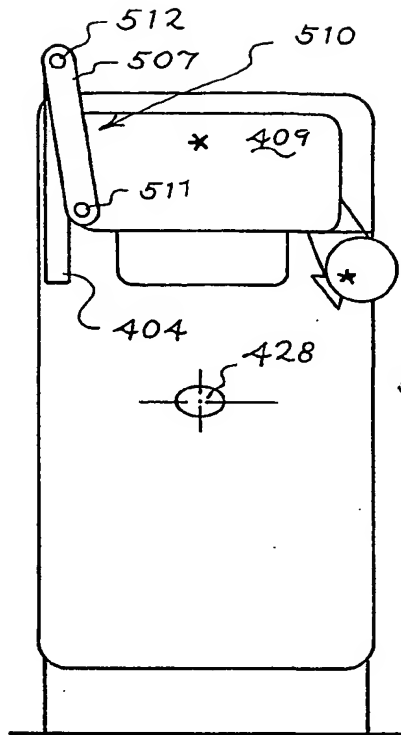


Fig. 20(a)

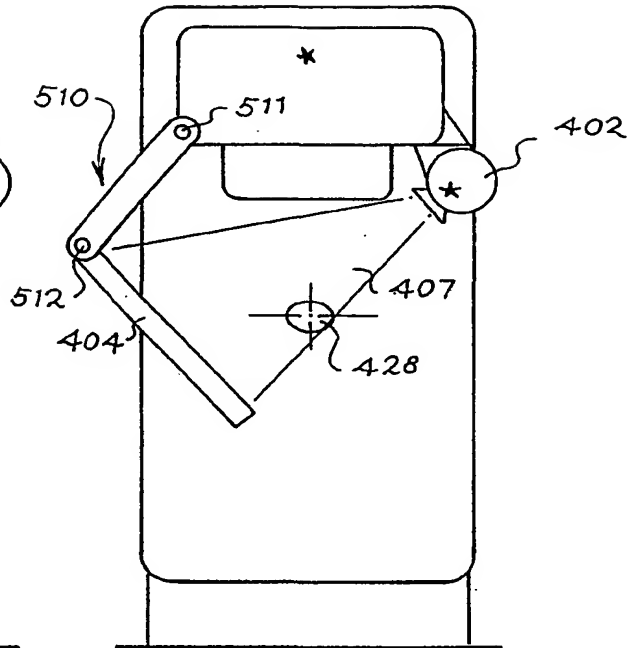


Fig. 20(b)

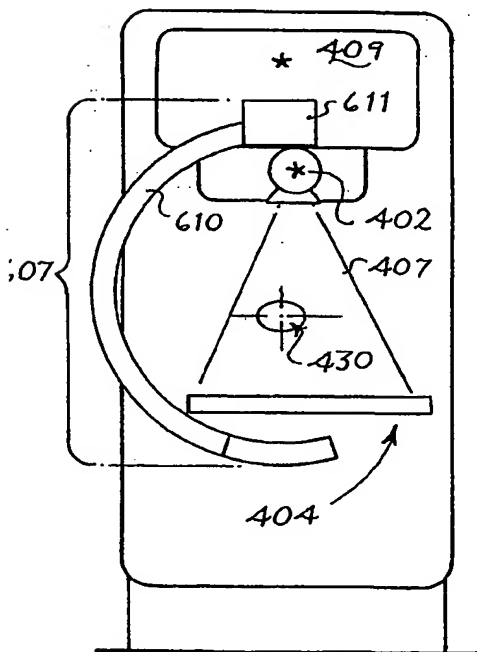


Fig. 21(a)

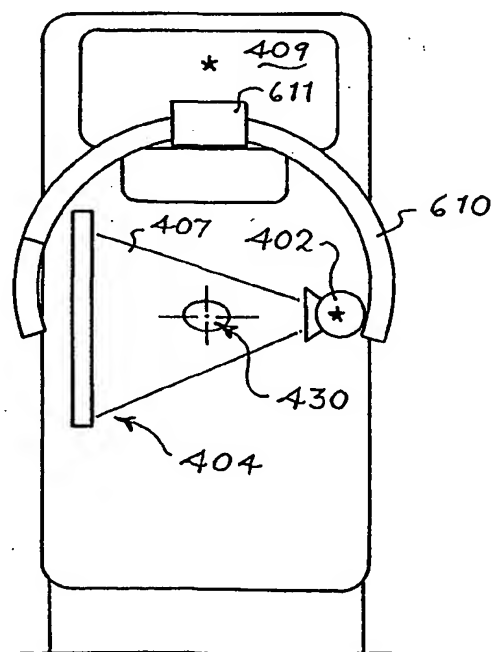


Fig. 21(b)

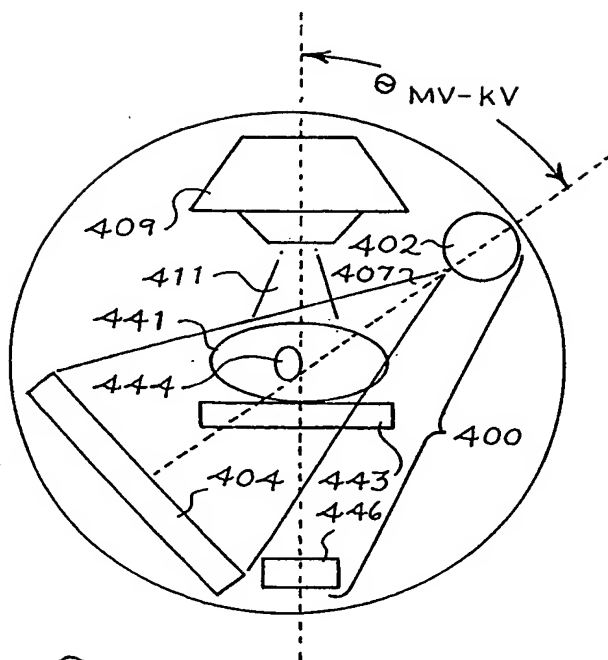


Fig. 23(a)

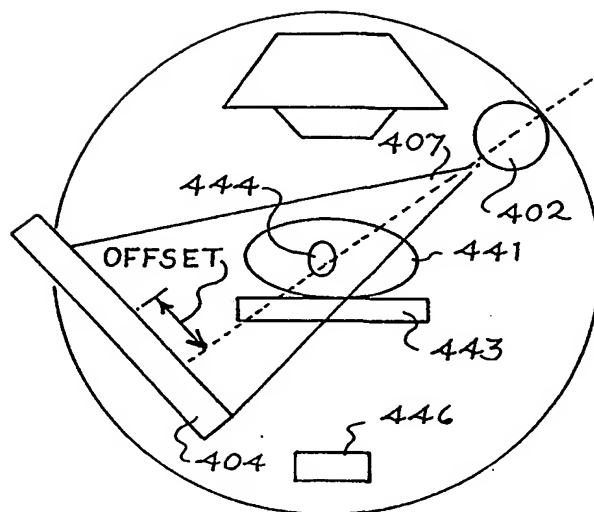


Fig. 23(b)

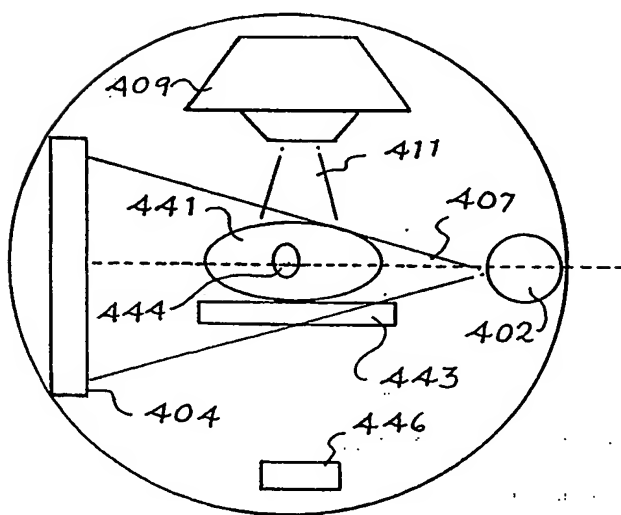


Fig. 23(c)

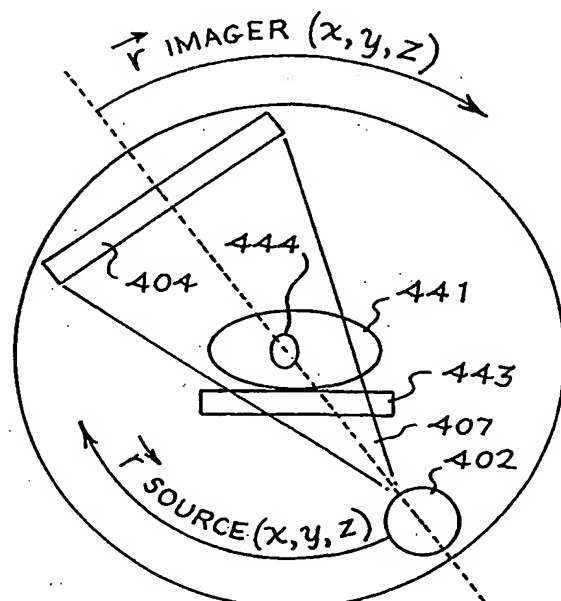


Fig. 23(d)

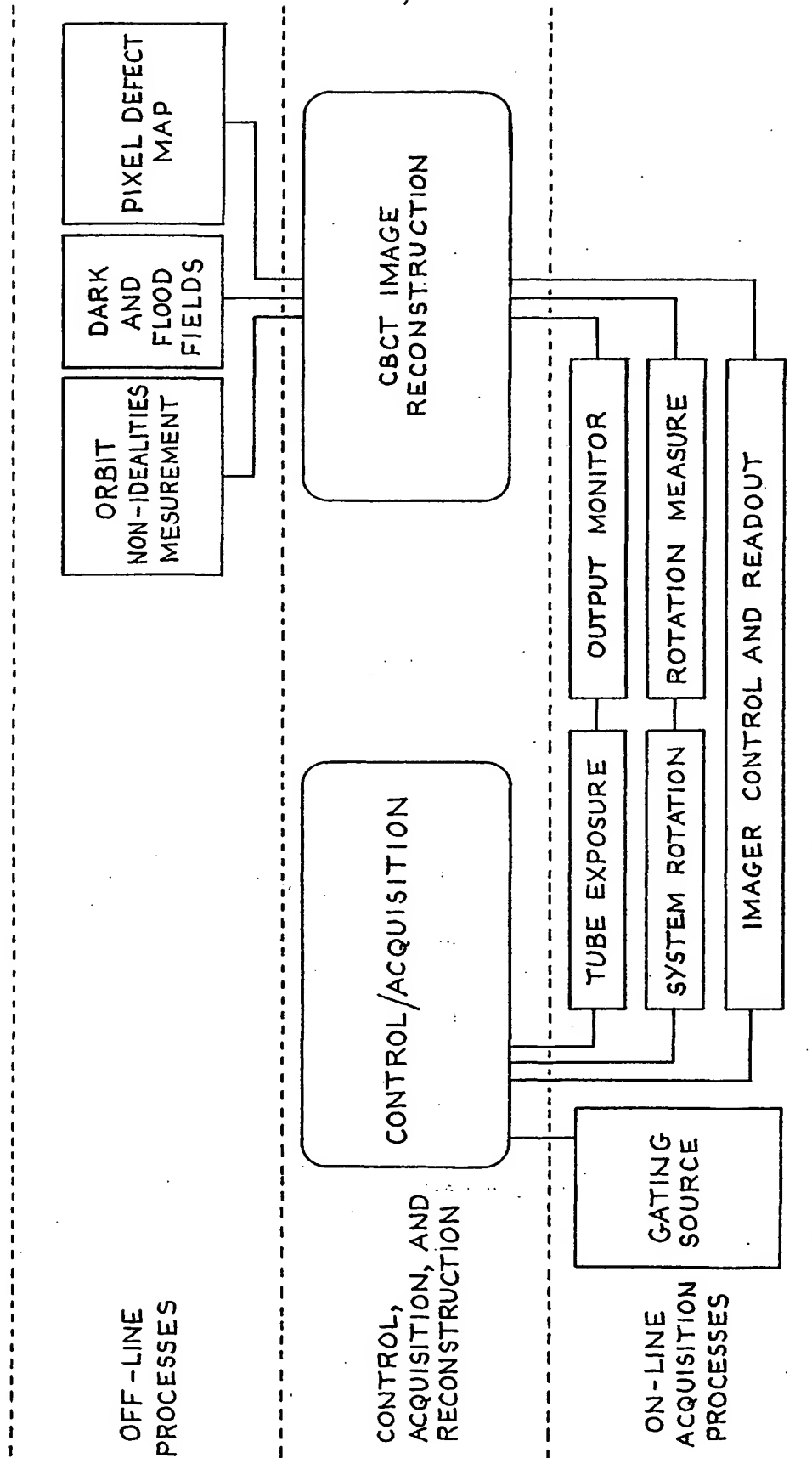
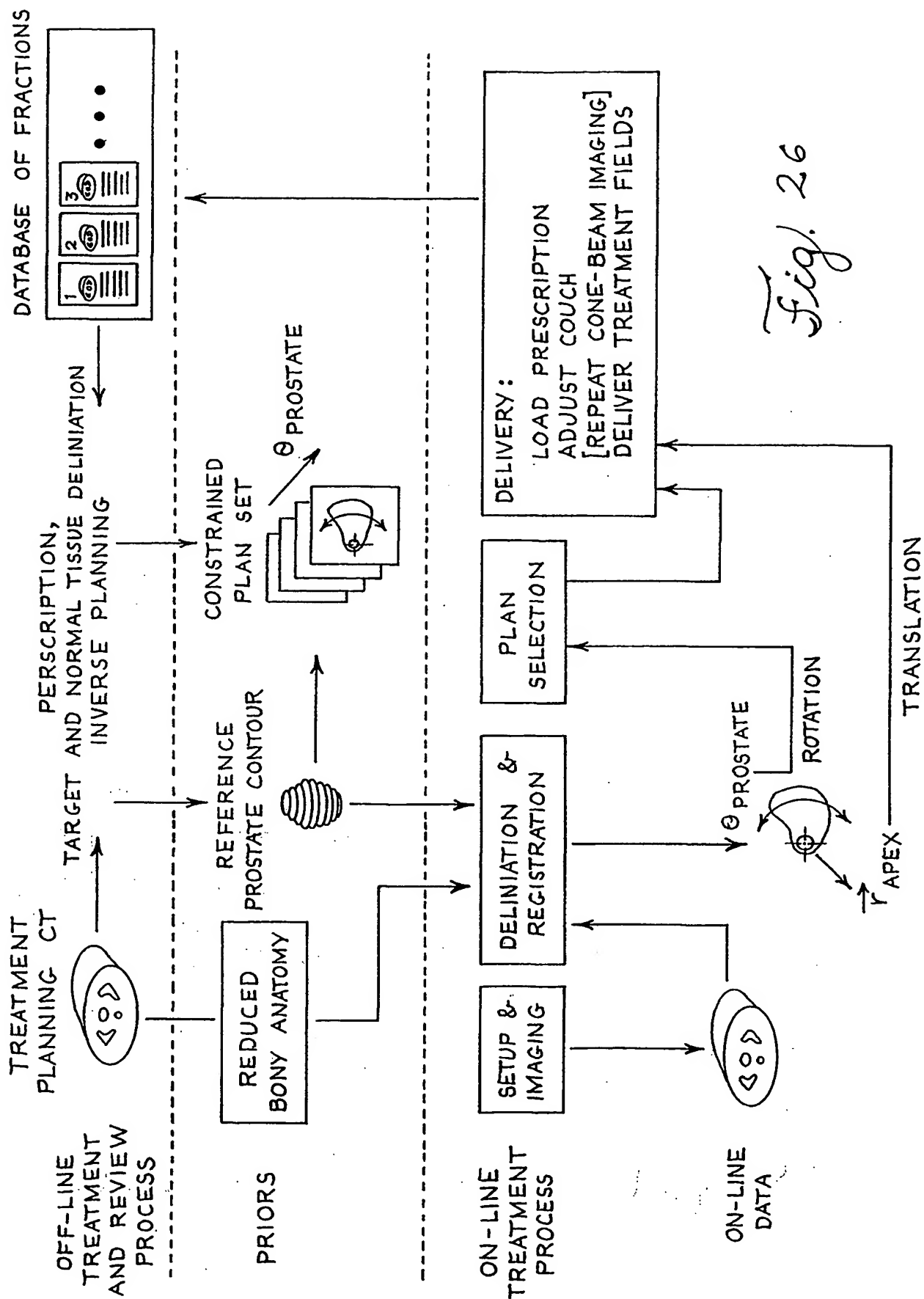


Fig. 24



(19) World Intellectual Property Organization
International Bureau



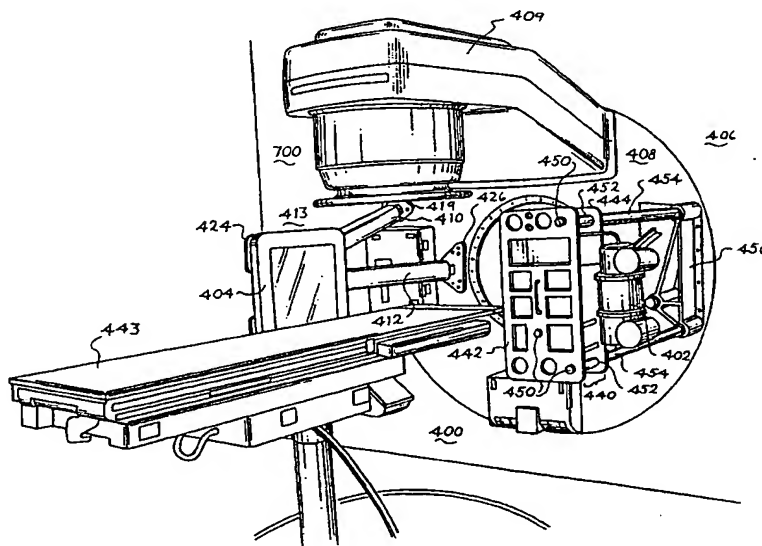
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PCT

(10) International Publication Number
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60/183,590 **18 February 2000 (18.02.2000)** **US**
- (71) Applicant: **WILLIAM BEAUMONT HOSPITAL**
[US/US]: **3601 West Thirteen Mile Road, Royal Oak, MI 48073-6769 (US).**
- (72) Inventors: **JAFFREY, David, A.:** 2476 Lincoln Road, Windsor, Ontario N8W 2R7 (CA). **WONG, John, W.:** 726 Tennyson Downs Court, Bloomfield Hills, MI 48304 (US). **SIEWERDSEN, Jeffrey, H.:** 2383 Timbercrest Court, Ann Arbor, MI 48105 (US).
- (74) Agent: **FREEMAN, John, C.:** Brinks Hofer Gilson & Lione, P.O. Box 10087, Chicago, IL 60610 (US).
- (81) Designated States (*national*): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW.
- (84) Designated States (*regional*): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).
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- (88) Date of publication of the international search report:
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- For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.*

(54) Title: **CONE-BEAM COMPUTERIZED TOMOGRAPHY WITH A FLAT-PANEL IMAGER**



(57) Abstract: A radiation therapy system that includes a radiation source (409) that moves about a path and directs a beam of radiation towards an object and a cone-beam computer tomography system (400). The cone-beam computer tomography system includes an x-ray source (402) that emits an x-ray beam in a cone-beam form towards an object to be imaged and an amorphous silicon flat-panel imager (404) receiving x-rays after they pass through the object, the imager providing an image of the object. A computer is connected to the radiation source and the cone beam computerized tomography system, wherein the computer receives the image of the object and based on the image sends a signal to the radiation source that controls the path of the radiation source.

WO 01/60236 A3

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US01/05077

A. CLASSIFICATION OF SUBJECT MATTER

IPC(7) : A61N 5/10

US CL : 378/65

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 378/65

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

NONE

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

EAST BRS: CT OR (compute\$5 tomograph\$2), amorphous silicon, cone-beam OR (cone beam).

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5,751,781 A (BROWN et al.) 12 May 1998 (12.05.1998), entire document.	81, 84, 86, 87

Y		1-35, 63-80
Y	US 5,748,700 A (SHEPHERD et al.) 05 May 1998 (05.05.1998), entire document.	1-35, 63-80
Y	US 5,661,773 A (SWERDLOFF et al.) 26 August 1997 (26.08.1997), entire document.	1-35, 63-80
A	US 5,912,943 A (DEUCHER et al.) 15 June 1999 (15.06.1999), entire document.	1-48, 63-88
A	US 6,152,598 A (TOMISAKI et al.) 28 November 2000 (28.11.2000), entire document.	36-48

☐

Further documents are listed in the continuation of Box C.

☐

See patent family annex.

Special categories of cited documents:	
"A" document defining the general state of the art which is not considered to be of particular relevance	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"E" earlier application or patent published on or after the international filing date	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"O" document referring to an oral disclosure, use, exhibition or other means	"&" document member of the same patent family
"P" document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

06 August 2001 (06.08.2001)

Date of mailing of the international search report

27 AUG 2001

Name and mailing address of the ISA/US

Commissioner of Patents and Trademarks
Box PCT
Washington, D.C. 20231

Facsimile No. (703)305-3230

Authorized officer

Allen C. Ho

Telephone No. (703) 308-6189

Form PCT/ISA/210 (second sheet) (July 1998)

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US01/05077

Box I Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)

This international report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claim Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☐ Claim Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claim Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of Item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:
Please See Continuation Sheet

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☒ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.: 1-48 and 63-88
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest ☐ The additional search fees were accompanied by the applicant's protest.
☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US01/05077

BOX II. OBSERVATIONS WHERE UNITY OF INVENTION IS LACKING

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1. In order for all inventions to be examined, the appropriate additional examination fees must be paid.

Group I, claim(s) 1-35, drawn to a radiation therapy system.

Group II, claim(s) 36-48, drawn to an x-ray imaging system.

Group III, claim(s) 49-55, drawn to an x-ray imaging system.

Group IV, claim(s) 56-62, drawn to an x-ray imaging system.

Group V, claim(s) 63-80, drawn to a method of treating an object with radiation.

Group VI, claim(s) 81-88, drawn to a method of adding an auxiliary imaging system to an existing radiation therapy system.

Group VII, claim(s) 89-91, drawn to a method of delineating a target volume located within a body and shown in a computerized image.

Group VIII, claim(s) 92, drawn to a method of delineating a target volume located within a body and shown in a computerized tomography image.

Group IX, claim(s) 93, drawn to a method of delineating a target volume located within a body and shown in a computerized tomography image.

The inventions listed as Groups I-IX do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons: The special technical feature of the Group I invention is the particular radiation therapy system. The special technical feature of the Group II invention is the particular x-ray imaging system. The special technical feature of the Group III invention is the particular x-ray imaging system. The special technical feature of the Group IV invention is the particular x-ray imaging system. The special technical feature of the Group V invention is the particular method of treating an object with radiation. The special technical feature of the Group VI invention is the particular method of adding an auxiliary imaging system to an existing radiation therapy system. The special technical feature of the Group VII invention is the particular method of delineating a target volume located within a body and shown in a computerized tomography image. The special technical feature of the Group VIII invention is the particular method of delineating a target volume located within a body and shown in a computerized tomography image. The special technical feature of the Group IX invention is the particular method of delineating a target volume located within a body and shown in a computerized tomography image. Since no common special technical feature exists between the inventions, unity of invention is lacking.